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Abd32888 Human can
Add13739 Osteoarth
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cytostatic; gene therapy; diagnostic marker; morbid state; osteoporosis;
neurological disease; Alzheimer's disease; Parkinson's disease; dementia;
      21-JAN-2003; 2003JP-00102206
09-MAY-2003; 2003JP-00131392
                                                                                                                                                                                                                                                                                                                                                                                                                                                                ADQ62941 standard; cDNA; 3122
                                                                21-JAN-2004; 2004EP-00001196
                                                                                                         28-JUL-2004.
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Add36459 Human aut
Ab164403 Stomach cc
Ad16403 Stomach cc
Ad113748 Osteoarth
Aca61195 Novel human
Ad040593 Human kin
Ad040593 Human kin
Ad040593 Human kin
Ad040593 Human gen
Acn44350 Human gen
Acn44350 Human gen
Acn44354 Human can
Aak65129 Human imm
Aak65129 Human ova
Ad167899 Human ova
Ad174276 Human ova
Ad174278 Human ova
Ad1374274 Human ova
Ad1374274 Human ova
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Ad139513 Human ova
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Ad139513 Human con
Acn19882 Breast ca
Ac158872 Human con
Aah10527 Human con
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Matches 3122;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Isogai T,
Yamamoto
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Claim
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                                          |GGGAGCCACCGTGGAGGCCAGGGCGGTGCAGAGACACGACGTGTGACTCGG
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Nagai K, Irie R;
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This invention relates to novel, isolated full length human cDNA molecules and the encoded proteins thereof. Specifically, it refers to cDNA clones obtained by an oligo-capping method, where none of these clones are identical to any known human mRNAs. The present invention describes an immunoassay to identify agonists and antagonists, as well antibodies, antisense molecules and siRNAs that can all be used to bind to and modulate expression of the cDNA molecules. As such, these molecules are useful for diagnostic markers or theraparities.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            gene; 88; human; oligo-capping method; diagnostic marker; gene therapy osteoporosis; neurological disease; Alzheimer's disease; parkinson's disease; dementia; short memory; cancer; sense or motor function; emotional reaction; fear response; panic; osteopathic; neuroprotective; nootropic; antiparkinsonian; cytostatic; osteopathic; neuroprotective; nootropic; antiparkinsonian; cytostatic;
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               GECGTEGCEGCECECECTEAGCACCCECAGCCTECEGCTCEAGGCEGAGGEGCEACTTC
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                                                                                                                              CC AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic CC activity, and can be used in gene therapy and vaccine production. (I) CC proteins and polymucleotides may be used in the prevention, diagnosis and CC treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat disorders associated with decreased CC expression by rectifying mutations or deletions in a patient's genome CC that affect the activity of (I) by expressing inactive proteins or to CC supplement the patients own production of (I). Additionally, (I) CC concers may be used to produce the secreted (I), by inserting the CC mucleic acids into a host cell and culturing the cell to express the CC protein. (I) proteins and polymucleotides may be used to prevent, CC cancers and cancer metastases of haematopoietic derived cells. AAK4703 CC cancers and cancer metastases of haematopoietic acityen genomic concers from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention.
Query Match
Best Local Similarity
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O7-NOV-2001 (first entry) Human immune/haematopoietic; immune/haematcycietic; immune/haema	
sequence SEQ ID NO:38235. opoietic antigen; cancer; s; ds.	
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Best Local
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08-DEC-2000; 2000US-0251869P.
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11-DEC-2000; 2000US-0251990P.
11-DEC-2000; 2000US-0254997P.
05-JAN-2001; 2001US-0259678P.
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19-MAY-2000;
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20-CCT-2000;
20-NOV-2000;
21-NOV-2000;
21-NO
AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis an treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I)
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                                                                                     The invention relates to isolated polynucleotide (I) and polypeptide (II) C sequences. (I) is useful as hybridisation probes, polymerase chain C reaction (PCR) primers, oligomers, and for chromosome and gene mapping, C and in recombinant production of (II). The polynucleotides are also used (in diagnostics as expressed sequence tags for identifying expressed (genes. (I) is useful in gene therapy techniques to restore normal (II) activity of (II) or to treat disease states involving (II). (II) is C useful for generating antibodies against it, detecting or quantitating a C useful for generating antibodies against it, detecting or quantitating a C supplement. (II) and its binding partners are useful in medical imaging C involving aberrant protein expression or biological activity. The C diagnostice and polynucleotide sequences have applications in C responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and C amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic patent did not appear in the printed specification, but was obtained in C electronic format directly from WIPO at
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             S
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23-AUG-2000;
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                                                                 ftp.wipo.int/pub/published_pct_sequences
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Best Local Similarity

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RESULT 7
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Human; probe; ss; gene expression; single exon probe; microarray; alternative splicing event; genomic alteration.
                                                                                                                                                                                                                        Human
US2003194704-A1.
                                                                                                                                                                                                                 genome derived single exon probe #20699
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(PENN/) (RANK/) (HANZ/) 03-APR-2002; 2002US-00029386 03-APR-2002; 2002US-00029386 PENN S G.
RANK D R.
HANZEL D K. Rank DR,

New human genome-derived single exon nucleic acid probes useful for human gene expression analysis, for identifying or characterizing alternative splicing events, for assessing genomic alterations or as tools for surveying tissues.

Claim 1; SEQ ID NO 20699; 80pp; English.

CC sequences in the specification, or their complements or fragments, and CC fully defined in the specification, or the 688 amino acid sequences (CC fully defined in the specification. The probe is a single exon probe that (CC hybridiese under high stringency conditions to a nucleic acid molecule (CC expressed in human cells or tissues. Also included are a spatially-CC addressable set of single exon nucleic acid probes for measuring human (CC purobes cited above, where each of the plurality of probes is separately (CC and addressably isolatable or amplifiable from the plurality), a single exon microarray for measuring human gene expression, a method of contiguous amino acids of any of the above-mentioned amino acid (CC sequences (optionally with conservative amino acid substitutions), an (CC sequences (optionally with conservative amino acid substitutions), an (CC acustomer desiring to measure gene expression, a method of contiguous amino acids of any of the above-mentioned amino acid storage medium which contains a database having a plurality of records (each record including data on the expression, a method of probes in cluding data by subscription, and a computer-readable compression analysis. The probes may be usef as tools for surveying contentive splicing events, in detecting and characterising contentive splicing events, in detecting and characterising gross alternative splicing events, in priming the synthesis of nucleic acids, or in expressing the ORF-encoded peptide. The present sequence is a human (CC patent did not form part of the princed apecification, but was obtained con alternation format directly from USPTO at (CC sequence data for this expression format directly from USPTO at (CC sequence data for this content and content of the propose are used in the sequence data for this content and content of the propose are used in the sequence data for this content and content of the propose are used content of the sequence and content of the invention. Note: The sequence data for this content of the c The invention relates to a nucleic acid probe for measuring human expression, comprising any of the 27,400 fully defined nucleotide gene

Sequence BP; 104 A; 279 C; 231 G; 94 T; 0 U; 0 Other;

Query Match
Best Local Similarity
Matches 707; Conserv Conservative 21.0**%**; 99.9**%**; <u>.</u> Score 657; Pred. No. 2. Mismatches DB 12; 2.9e-296; Length 708 Indels <u>,</u> Gaps

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RESULT 8
ACH73793/c
ID ACH73793 standard;
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                                                                                                 US2003194704-A1
                                                                                                                                          Human; probe; ss; gene expression; single exon probe; microarray; alternative splicing event; genomic alteration.
                                                                                                                                                                            Human genome derived single exon probe #6988.
                                                                                                                                                                                                   29-JUL-2004
(PENN/)
                                03-APR-2002; 2002US-00029386
                                                     03-APR-2002; 2002US-00029386
                                                                           16-OCT-2003
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New human genome-derived single exon nucleic acid probes useful for human gene expression analysis, for identifying or characterizing alternative splicing events, for assessing genomic alterations or as tools for
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                                                                                                                                                                                                                       Rank DR,
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Claim 15; SEQ ID NO 6988; 80pp; English.

CC measuring human gene expression, a vector comprising the single exon CC probe cited above, an ORF-encoded peptide comprising at least 8 CC contiguous amino acids of any of the above-mentioned amino acid CC sequences (optionally with conservative amino acid substitutions), an CC methods of selling and/or licensing single exon probes or microarrays to CC methods of selling and/or licensing single exon probes or microarrays to CC methods of selling and/or licensing single exon probes or microarrays to CC methods of selling to measure gene expression, a method of providing CC human gene expression data by subscription, and a computer-readable storage medium which contains a database having a plurality of records (each record including data on the expression of a single exon probe CC cited above. The probe, methods and apparatus are useful in gene CC expression analysis. The probes may be used as tools for surveying tissues to detect the presence of expressed messages that contain their CC specific exon, or in constructing genome-derived single exon microarrays. CC In addition, the probes are used in identifying and characterising gross calternative splicing events, in detecting and characterising gross calternative splicing events, in detecting and characterising gross calternative splicing the ORF-encoded peptide. The present sequence is a human single exon probe of the invention. Note: The sequence is a human considerated in electronic format directly from USFTO at sequence. html?DocID=20030194704 The invention relates to a nucleic acid probe for measuring human gene expression, comprising any of the 27,400 fully defined nucleotide sequences in the specification, or their complements or fragments, and encoding at least 8 amino acids of any of the 6888 amino acid sequences fully defined in the specification. The probe is a single exon probe that hybridises under high stringency conditions to a nucleic acid molecule expressed in human cells or tissues. Also included are a spatially-addressable set of single exon nucleic acid probes for measuring human gene expression (comprising a plurality of single exon nucleic acid probes cited above, where each of the plurality of probes is separately and addressably isolatable or amplifiable from the plurality), a single exon microarray for measuring human gene expression, a method of

Sequence 524 BP; 61 A; 212 C; 162 G; 89 T; 0 U; 0 Other;

Similarity

16.8%; Score 524; DB 12; 100.0%; Pred. No. 4.3e-234;

Length 524;

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Best Local Simi
Matches 524;
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CCGAGCGAGTCACGGACCATGAAGAGCGTTCGTGCCGCGGCCCAAGGCCGGGATGGGG
                                                                      CCGTCCGCAGAGGCGCACGTCGAGGGTCCCCGGGCGCTCCCGTGGACGTTGGCGGTAGCG
                                                                                                             GCGCCTGGGGAGGATGGACGAGGGAGCGGGGGACCGCTAACGGGGCTCCCTCTGCGCGCC
                                                                                                                                                                                                          TTCGCAGGGAGCCACCGTGGAGGCCAGGGCGGTGCAGAGACACGACGTGTGACTCGGAGT
                                                  CCGTCCGCAGAGGCGCACGTCGAGGGTCCCGGGGCGCGCTCCGTGGACGTTGGCGGTAGCG
                                                                                                                                            GCGCCTGGGGAGGATGGACGAGGGAGCGGGGGACCGCTAACGGGGCTCCCTCTGCGCGCCC
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11 3 2 2 2 2 3 3 3 3 3 3 5 5 5 5 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	Db D
\$2785 standard; cDNA; 973 BP \$2785; \$400V-2001 (first entry) \$100V-2001 (first entry) \$100V-2	284 C 784 G 224 G 224 G 844 C 164 C 904 C 104 C 104 C 44 T
igen encoding cDNA SEQ ID NO:7845. immune/haematopoietic antigen; cancer; ne; metastasis; ss.	CCGAGCGAGTCACCGACCATGAAGAGCGTTCGTGCCGCGCGCG
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2000US-0232399P. 2000US-023401P. 2000US-023401P. 2000US-023401P. 2000US-023405P. 2000US-023465P. 2000US-0234997P. 2000US-0234997P. 2000US-0235834P. 2000US-0235834P. 2000US-0235657P. 2000US-023657P. 2000US-024653P. 2000US-0246611P.	022718; 0228902 0228928; 022928; 022934; 022934; 022950; 0229510; 0229513; 023043; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124; 023124;

17-NOV-2000; 17-NOV-2000; 17-NOV-2000;

-NOV-2000;

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                                                                                                                                                                                       CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic cattivity, and can be used in gene therapy and vaccine production. (I) CC cattivity, and can be used in gene therapy and vaccine production. (I) CC proteins and polynucleotides may be used in the prevention, diagnosis and CC treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat disorders associated with decreased CC expression by rectifying mutations or deletions in a patient's genome CC that affect the activity of (I) by expressing inactive proteins or to CC supplement the patients own production of (I). Additionally, (I) CC polynucleotides may be used to produce the secreted (I), by inserting the CC protein. (I) proteins and polynucleotides may be used to prevent, CC diagnose and treat immune/haematopoietic-derived cells. AAK4703 CC cancers and cancer metastases of haematopoietic acitigen genomic CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169 cc represent sequences used in the exemplification of the present invention.
                                                                                                     Query Match
Best Local Simi
Matches 624;
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17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
01-DEC-2000;
05-DEC-2000;
05-DEC-2000;
05-DEC-2000;
06-DEC-2000;
06-DEC-2000;
08-DEC-2000;
08-DEC-2000;
18-DEC-2000;
11-DEC-2000;
11-DEC-2000;
                                                                                                                                                                 Sequence 973 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Claim 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       (HUMA-) HUMAN GENOME SCI INC
                                                                                                                      Local Similarity
                  2018
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TTCCCGGACAAGAAAATTGCAATCAAATGTCAGCAGCTTTTATTACCTTAATCTTTCAG
                                                                                                                                                                                                                                                                                                                                                                                                                                                                   SEQ ID NO 7845; 3071pp + Sequence Listing; English.
                                              Barash SC,
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2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249218P.
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2000US-0249264P.
2000US-0249265P.
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2000US-0249299P.
2000US-0249299P.
2000US-025999P.
2000US-025999P.
2000US-025190P.
2000US-0251869P.
2000US-0251990P.
                                                                                                         Conservative
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                                                                                                                                                                 255 A; 253 C; 226 G; 234 T; 0 U; 5 Other;
                                                                                                                    16.1%;
                                                                                                     Score 504; DB 4; L
Pred. No. 9.3e-225;
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RESULT 10
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ID AAK834
AC AAK834
AC AAK834
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29-JUN-2000
30-JUN-2000
37-JUN-2000
07-JUL-2000
07-JUL-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Human; immune; haematopoietic; immune/haematopoietic cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                            31-JAN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human immune/haematopoietic antigen genomic sequence
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AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis an treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the
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useful f
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AAK83427
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                                                                                                                                             09-AUG-2001.
                                                                                                                                                                                                                                        cytostatic;
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09-NOV-2000

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01-DEC-2000

08-DEC-2000

08-DEC-2000
AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis an
                                                                                                                Nucleic
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2000US-0241785P.
2000US-0241786P.
2000US-0241787P.
2000US-0241809P.
2000US-0246476P.
2000US-0246475P.
2000US-0246475P.
2000US-0246478P.
2000US-0246478P.
2000US-0246524P.
2000US-0246528P.
2000US-0246510P.
2000US-0246510P.
2000US-0246510P.
2000US-0246510P.
2000US-0246510P.
2000US-0249210P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-02492189P.
2000US-0251856P.
2000US-0251856P.
2000US-0251869P.
2000US-0251989P.
                                                                     SEQ
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                                                                                                                                                                    SC,
                                                                    NO 38239; 3071pp +
                                                                                                                                                                                                 SCI
                                                                                              human immune/hematopoietic diagnosing and/or treating
                                                                                                                                                                     Ruben
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                                                                    Sequence
                                                                    Listing;
prevention, diagnosis and
                                                                                                antigen polypeptides, cancers and metastasis.
                                                                    English
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Best Local S
Matches 471
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Sequence 476 BP; 59 A; 190 C; 157 G; 70 T; 0 U; 0 Other;
                              1136
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               GCGACGCTCGCTGGAGCTGGGCGCCGCCGCTTCCCCGCTGCACGCCGCCGCGGCG
                                                                                                                                                          GCAGAACCTGCGGCAGGAGCTGCAAAAGACGCGCCAGAAGGCGCAGGAGCTGGCGGTGTC
                                                                                                                                                                                                                          GCAGAACCTGCGGCAGGAGCTGCCAAAAGACGCGCCAGAAGGCGCGCGAGGAGCTGGCGGTGTC
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                                                                                                                                                                                                                                                                                                                                         GCCCAGCCGAGCCCACCGCGATGGCGAGGGAGTGCAAGGCGCTGCTGGACGGCTC
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GCGACGCTCGCTGGAGCTGGGCGCCGCGTTCCCGCTGCACGCGCCGCGGCG
                                                               CGAGTTCGAGCGGCTCTGGGTGGCCTTCTCGGGCTGCCTGGACCTGCTGGAAGCGGACAT
                                                                                   CGAGTTCGAGCGGCTCTGGGTGGCCTTCTCGGGCTGCCTGGACCTGCTGGAAGCGGACAT
                                                                                                                                   CAACAAGACGACTGCGTGCTACCACCACCTGGTGCTGACCGTCGGTGGCTCGGCGGACTC
                                                                                                                                                                                                                                                                                                                                                                         GGCCCAGCCGGAGCCCACCGCGATGGCGAGGGAGGAGTGCAAGGCGCTGCTGGACGGGCT
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ilarity 100.0%;
Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              0,
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Pred. No. 2.5e-209;
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RESULT 12
AAK83426/c
07-NOV-2001
                                                                                                                                                   AAK83426
                 09-AUG-2001.
                                                                             Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                              Human immune/haematopoietic antigen genomic sequence
17-JAN-2001; 2001WO-US001354.
                                   WO200157182-A2
                                                                    cytostatic;
                                                                                                                                                   standard;
                                                                     gene
                                                                                                                 (first entry)
                                                                      therapy; vaccine; metastasis;
                                                                                                                                                   DNA;
                                                                                                                                                    ₿₽
                                                                                                SEQ
                                                                                                ID NO:38238.
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2000; 2000U 2000; 2000U 2000; 2000U 2000; 2000U 2000; 2000U 2000; 2000U	2-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 4-SEP-2000; 2000U 1-SEP-2000; 2000U 1-SEP-2000; 2000U 1-SEP-2000; 2000U 1-SEP-2000; 2000U 1-SEP-2000; 2000U 1-SEP-2000; 2000U		28-UN-2000; 2000US-0214886F. 30-UN-2000; 2000US-0214886F. 30-UN-2000; 2000US-0216880P. 07-UL-2000; 2000US-0216880P. 11-UL-2000; 2000US-0217487P. 11-UL-2000; 2000US-0217496P. 11-UL-2000; 2000US-0217496P. 11-UL-2000; 2000US-022964P. 14-UL-2000; 2000US-0229519P. 14-AUG-2000; 2000US-0224518P. 14-AUG-2000; 2000US-0225213P. 14-AUG-2000; 2000US-0225213P. 14-AUG-2000; 2000US-0225214P. 14-AUG-2000; 2000US-0225214P. 14-AUG-2000; 2000US-0225214P. 14-AUG-2000; 2000US-0225216P. 14-AUG-2000; 2000US-022526FP. 14-AUG-2000; 2000US-022526P. 14-AUG-2000; 2000US-022526P. 14-AUG-2000; 2000US-022575PP.	1-JAN-2000; 2000U 4-FEB-2000; 2000U 4-FEB-2000; 2000U 6-MAR-2000; 2000U 6-MAR-2000; 2000U 7-MAR-2000; 2000U 8-APR-2000; 2000U 8-APR-2000; 2000U
PA (HUMA-) HUMAN GENOME SCI INC. XX PI Rosen CA, Barash SC, Ruben SM; XX DR WPI; 2001-483426/52. PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides, PT useful for preventing, diagnosing and/or treating cancers and metastasis	01-DEC-2000 05-DEC-2000 05-DEC-2000 05-DEC-2000 05-DEC-2000 06-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000 08-DEC-2000	17-NOV-2000 17-NOV-2000	PR 20-CCT-2000; 2000US-02417859. PR 20-CCT-2000; 2000US-0241787P. PR 20-CCT-2000; 2000US-0241808P. PR 20-CCT-2000; 2000US-0241808P. PR 20-CCT-2000; 2000US-0241808P. PR 20-CCT-2000; 2000US-0241826P. PR 20-CCT-2000; 2000US-0244617P. PR 08-NOV-2000; 2000US-0246474P. PR 08-NOV-2000; 2000US-0246475P. PR 08-NOV-2000; 2000US-0246477P. PR 08-NOV-2000; 2000US-0246477P. PR 08-NOV-2000; 2000US-0246478P. PR 08-NOV-2000; 2000US-0246523P. PR 08-NOV-2000; 2000US-0246528P. PR 08-NOV-2000; 2000US-0246528P. PR 08-NOV-2000; 2000US-0246528P. PR 08-NOV-2000; 2000US-024653P. PR 08-NOV-2000; 2000US-024653P. PR 08-NOV-2000; 2000US-024653P. PR 08-NOV-2000; 2000US-024653P. PR 08-NOV-2000; 2000US-024651P. PR 08-NOV-2000; 2000US-024651P. PR 08-NOV-2000; 2000US-024661P. PR 08-NOV-2000; 2000US-024661P. PR 08-NOV-2000; 2000US-024661P.	02-0CT-2000, 02-0CT-2000, 02-0CT-2000, 02-0CT-2000, 02-0CT-2000, 13-0CT-2000, 13-0CT-2000, 20-0CT-2000,

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ID AAS725
XX AAS725
AC AAS725
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DT 13-FEB
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DE DNA en
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Query Match
Best Local Similarity
Matches 427; Conserv
                Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Sequence 476 BP; 58 A; 191 C; 157 G; 70 T; 0 U; 0 Other;
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                                                                    DNA encoding novel human diagnostic protein #8312.
                                                                                                         13-FEB-2002
                                                                                                                                              AAS72508
                                                                                                                                                                             AAS72508 standard; cDNA; 1349
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            SEQ ID NO 38238; 3071pp + Sequence Listing; English
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                                                                                                           (first
                                                                                                                                                                                                                                                                      50
                                                                                                           entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Score 427; DB 4; L; pred. No. 9.3e-189; 0; Mismatches 0;
                                                                                                                                                                               ВP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         The invention relates to isolated polynucleotide (I) and polypeptide (II) CG sequences. (I) is useful as hybridisation probes, polymerase chain CG reaction (PCR) primers, oligomers, and for chromosome and gene mapping, CG and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal CG useful for generating antibodies against it, detecting or quantitating a CG useful for generating antibodies against it, detecting or quantitating a CG useful for generating antibodies against it, detecting or quantitating a CG supplement. (II) and its binding partners are useful in medical imaging CG supplement (II) and its binding partners are useful for treating disorders involving aberrant protein expression or biological activity. The CG diagnostics, forensics, gene mapping, identification of mutations CG diagnostics, forensics, gene mapping, identification of mutations CG diagnostics, forensics gene mapping, identification of mutations CG amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention. Note: The sequence data for this control of mutations in the printed specification, but was obtained in CG electronic format directly from WIPO at the control of mutations in the printed specification, but was obtained in CG electronic format directly from WIPO at the control of the
                                                                                                                                                                                                                                                                                                                                                                                                                                                          Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Query Match
Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Sequence 1349 BP; 171 A; 472 C; 483 G; 223 T;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    New isolated polynucleotide and encoded polypeptides, useful diagnostics, forensics, gene mapping, identification of mutat responsible for genetic disorders or other traits and to assebiodiversity.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        31-MAR-2000; 2000US-00540217.
23-AUG-2000; 2000US-00649167.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      WPI; 2001-639362/73.
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                                                                                                                                                            282
                                                                                                                                                                                                                                                                                                                                                              402
                                                                                                                                                                                                                                                                                                                                                                                                                                                          241;
                                                                                                                                                                                                                                                                                                                                                                                   GACGACTGCGTGCTACCACCACCTGGTGCTGACCGTCGGTGGCTCGGCGGACTCGCAGAA
            C 1142
                                                                                            CGAGCGGCTCTGGGTGGCCTTCTCGGGGCTGCCTGGACGTGCTGGAAGCGGAACATGCGACG 1141
                                                                                                                                                            CGCCCGGCTGACTGCTGCTGCGCGCGACCGGGGCCTGGCCGCCGACGAGCGCGAGTT
                                                                                                                                                                                                                                                          CCTGCGGCAGGAGCTGCAAAAGACGCGCCAGAAGGCGCAGGAGCTGGCGGTGTCCACCTG
                                                                                                                                                                                                                                                                                                        CCTGCGGCAGGAGCTGCAAAAGACGCGCCAGAAGGCGCAGGAGCTGGCGGTGTCCACCTG
                                                                                                                                                                                                                                                                                                                                                            GACGACTGCGTGCTACCACCACCTGGTGCTGACCGTCGGTGGCTCGGCGGACTCGCAGAA
                                                             CGAGCGGCTCTGGGTGGCCTTCTCGGGCTGCCTGGACCTGCTGGAAGCGGACATGCGACG
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RESULT 14
ABN50582
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                                                                                                                                                                                                                                                                               The present invention describes oligonucleotide libraries for detecting messenger RNAs that populate a [sub-)transcriptome, where the (sub-)transcriptome comprises messenger RNAs transcribed from multiple conting that populate a genome. The library comprises several coligonucleotides, each capable of hybridising selectively to a set of coligonucleotide per RNAs transcribed from a given transcripton unit of the genome, which encodes one or more messenger RNA splice variants. The coligonucleotide libraries are useful for detecting mRNAs from a coligonucleotide libraries are useful for detecting mRNAs from a coligonucleotide libraries are useful for detecting mRNAs from a coligonucleotide libraries are useful for detecting mRNAs from a coligonucleotide libraries are useful for detecting mRNAs from a coligonucleotide libraries may also be used as specialised minicular colibraries to detect transcripts of a sub-transcriptome under a particular biological or pathological state, and so allowing the detection of tissue coligonucleotide specific genes such as those genes only expressed in coligonucleotide tissue under a specific pathological condition; to detect coligonerate specific genes and to detect man transcripts and splice variants of a transcriptome of a pathent suffering from a particular colisorder. ABN27253 to ABN59589 represent oligonucleotide sequences from corresponding from a particular colores in the exemplification of the present invention. N. The semence data for this meter of the corresponding from a particular colores in the exemplification of the corresponding from a particular colores in the scent of the corresponding from a particular colores in the scent of the corresponding from a particular colores in the scent of the corresponding from a particular colores in the scent of the corresponding from a particular colores in the scent of the corresponding from a particular colores in the scent of the corresponding from a particular coloresponding from a particular colores in the scent of the correspo
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Best Local Similarity
                                                                                                                                                                             Sequence
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription unit genome, useful for detecting tissue-, pathology-, and developmental-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         ABN50582 standard; DNA; 60
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Example 1; SEQ ID NO 23330; 47pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  specific
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    splice variant; transcriptome; oligonucleotide library; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Human spliced transcript detection oligonucleotide SEQ ID NO:23330.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     15-JUL-2002
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Shoshan A, Wasserman A,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   28-JUL-2000; 2000US-0221607P
02-MAY-2001; 2001US-0287724P
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                                                                                                                                                                                                                                                                          present
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           (COMP-) COMPUGEN INC.
                            of the printed specification,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            mouse; rat; splice transcript; detection;
                                                                                                                                                                                                                     invention. N.B. The sequence data for this patent did not the printed specification, but was obtained in electronic y from WIPO at ftp.wipo.int/pub/published_pct_sequences
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  BP; 11 A; 14 C; 16 G; 19 T; 0 U; 0 Other;
                                                                                        Conservative
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Pred. No.
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RESULT 15
ABK83571/c
              CC (GCA), by detecting the level of expression of gene(s) (GC) and while analysis as given in the specification, and comparing the comparison level to an expression level in an unactivated GC, where CC expression of at less one of Gs is indicative of GCA. Also included are CC modulating (MC) GA by contacting GC with an agent that alters the CC expression of at less tone gene in Gs; (2) screening (M3) for an agent CC tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease using the gene expression CC pathogen or sterile inflammatory disease, by detecting the level of CC expression in a sample of the tissue of gene(s) from Gs, where the level of expression at reile inflammatory disease, by detecting the level of CC expression in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by detecting the level of CC expression in a subject, exposure of in a tissue, an allergic response in a subject to a pathogen or sterile conformation (especially chronic) in a tissue, an allergic response in a subject to a pathogen or sterile conformation of the gene is indicative of inflammation; (4) treating CC (MS) an inflammatory disease, by contacting a tissue having inflammation with an CC is useful for detecting GCA; M2 is useful for modulating GA; M3 is useful CC for screening an agent capable of modulating GCA preferably in an CC inflammation in a tissue; M1 is useful for detecting GA; M3 is useful for detecting GA; M3 is useful for detecting GA; M3 is useful for detecting GCA; M2 is useful for detecting GA; M3 is useful for detecting GCA; M3 is useful GC disease, ulcerative colitis, periodontal disease; also bacterial infection, viral infection, parasitic infection, protozoal infection, conditions. The present sequence represents a gene differentially conditions. The present sequence represents a gene differentially disease.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Detecting granulocyte activation by detecting differential expression of genes associated with granulocyte activation, which serves as diagnostic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human; 88; granulocytic cell; DNA chip; bacterial infection; viral infection; parasitic infection; protozoal infection; fungal infection; sterile inflammatory disease; psoriasis; rheumatoid arthritis; glomerulonephritis; asthma; thrombosis; cardiac reperfusion injury; renal reperfusion injury; ARDS; adult respiratory distress syndrome; inflammatory bowel disease; crohn's disease; ulcerative colitis; periodontal disease;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Claim 1; SEQ ID NO 142; 114pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Beazer-Barclay Y, Weissman SM,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    granulocyte activation; chronic inflammation; allergy
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             The invention relates to detecting (M1) granulocyte (GC) activation
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        markers that is useful for monitoring disease
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   03-OCT-2000; 2000US-0237189P
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ADI/13596;

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ADI/13596;

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O6-MAY-2004 (first entry)
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O3-JUL-2003
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O3-JUL-2003.
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O3-JUL-2003.
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O1-DEC-2002; 2002WO-US041225.
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O2-DEC-2002; 2002WO-US041225.
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O2-DEC-2001; 2001US-0342603P.
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O2-DEC-2001; 2001US-0342603P.
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O3-JUL-2003.
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O3-DEC-2001; 2001US-0342603P.
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O6-MAY-2004; 2001US-0342603P.
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O6-MAY-2004; 2002WO-US041225.
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OBC-2002; 2002WO-US041225.
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O7-DEC-2002; 2002WO-US041225.
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O7-DEC-2003; 2002WO-US041225.
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O7-DEC-2003; 2002WO-US041225.
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                                                                                                                                                                                   The invention relates to a method of determining susceptibility of an clinity individual to joint space marrowing and/or osteophyte development and/or joint pain comprising identifying whether the individual has at least one polymorphism in a polymucleotide encoding at least one of the protein listed in the specification. The methods, composition and agent are useful for modulating the susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain that is associated with a disease, preferably osteoarthritis. The cell line and the non-human animal are useful for screening for an agent for diagnosing an individual having susceptibility to joint space narrowing and/or osteophyte development and/or joint pain. This sequence corresponds to the polymucleotide encoding a protein listed in the specification. (Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format directly from WIPO at fip.wipo.int/pub/published_pct_sequences).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Query Match
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Query Match
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Matches 53; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              ds; gene; osteopathic; antiinflammatory; antiarthritic; gene therapy;
joint space narrowing; osteophyte development; joint pain;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Sequence 175737 BP; 41985 A; 43790 C; 42407 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Osteoarthritis-associated polymorphic nucleotide
                                                                                                                                   Sequence 175737 BP; 41985 A; 43790 C; 42407 G; 47555 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Disclosure; SEQ ID NO 128; 297pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         47829 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 47777
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   ilarity 100.0%; I Conservative 0;
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tive 0; Mismatches
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100.0%; Pred. No.
                                  1.7%; Score 53;
100.0%; Pred. No.
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                                                                     DB 10;
                               4.9e-14;
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                                                                 Length 175737;
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AAC03795 standard;

CDNA; 381

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RESULT 17
ADQ18934/c
ID ADQ189
XX ADQ189
XX ADQ189
XX ADQ189
XX ADQ189
XX Boft t
XX Soft t
XX Human
XX Homo s
XX Hom
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           닭
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                                                                                                                                 片
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RESULT 18
AAC03795/c
                                                                                                                                                                                                                                                                Query Match
Best Local S
Matches 53
                                                                                                                                                                                                                                                                                                                                                                                                                                                         which comprises obtaining a first soft tissue sample from an individual and a normal soft tissue sample from the same or different individual, determining the expression of a gene in both samples and comparing the expression of the gene in both soft tissue samples, where a higher level of protein expression in the first soft tissue sample indicates the presence of soft tissue sarcoma. The method of the invention has cytostatic applications and may be useful for detecting soft tissue sarcoma, possibly via gene therapy or vaccine production. The nucleic acid sequences may be useful in diagnostic and screening applications. The current sequence is that of a human soft tissue sarcoma-upregulated DNA of the invention. The current sequence is not shown within the specification per se but was submitted in CD format by the inventor.
                                                                                                                                                                                                                                                                                                                                                                                        Sequence 175737 BP; 41985 A; 43790 C; 42407 G; 47555 T; 0 U; 0
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Early detection of soft tissue sarcoma comprises determining expression of a gene in a first soft tissue sample and a normal soft tissue sample and comparing the gene expression, also useful in treating soft tissue
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Human soft tissue sarcoma-upregulated DNA - SEQ ID 1753.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Aziz N,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                26-NOV-2002; 2002US-0429739P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 26-NOV-2003; 2003WO-US038193
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The invention relates to a novel method for detecting soft tissue sarcoma
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              WPI; 2004-441208/41
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Example 2; SEQ ID NO 1753; 210pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               (PROT-)
                                                                                                                                                                                                                                                                                                  ocal Similarity
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                                                                                                                                                                       2888 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT
                                                                                                                                                                                                                                                                   53;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               PROTEIN DESIGN LABS INC.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 47777
                                                                                                                                     TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT
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                                                                                                                                                                                                                                                             1.7%; Sollarity 100.0%; I Conservative 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             sarcoma;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Zlotnik A;
                                                                                                                                                                                                                                                                                                      Score 53;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              ВP
                                                                                                                                                                                                                                                                       Mismatches
                                                                                                                                                                                                                                                                                                                                        DB 12;
                                                                                                                                                                                                                                                                                                  4.9e-14;
                                                                                                                                                                                                                                                                                                                                 Length 175737;
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                                                                                                                                        47777
                                                                                                                                                                                                                                                                                                                                                                                                         Other;
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RESULT 19
AAS78337/G
ID AAS78
XX AAS78
AC AAS78
XX IN 13-FE
XX UNN E
XX HUMAI
KW HOMO
XX HOMO
XX HOMO
XX WO20
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                                                                                                                                                                                                                                                                                                                                                                                     Matches 52;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   mRNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design expression and secretion
                                                                            Human; chromosome mapping; gene mapping; gene therapy; food supplement; medical imaging; diagnostic; genetic o
                                                                                                                             DNA encoding novel human diagnostic protein #14141.
                                                                                                                                                                13-FEB-2002
                                                                                                                                                                                                                                 AAS78337 standard; cDNA; 1437 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                      Sequence 381 BP; 73 A; 98 C; 84 G; 123 T; 0 U; 3 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      mRNAs encoding secreted proteins. An ORF has been identified within the sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         New nucleic acid that is a 5' expressed sequence tag (5' EST) for obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for diagnostic, forensic, gene therapy and chromosome mapping procedures.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Dumas Milne Edwards J,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     06-SEP-2000
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                 WO200175067-A2
                                                Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          The present sequence is one of a large number of 5' ESTs derived from
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Claim 1; SEQ ID NO 3793; 71pp + Sequence Listing; English
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Human; 5' EST;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Human secreted
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         06-OCT-2000
                                                                                                                                                                                                                                                                                                                                    3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAGAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               2000-500381/45
DB; AAG03789.
                                                                                                                                                                                                                                                                                                                     107
                                                                                                                                                                                                                                                                                                                   CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 56
                                                                                                                                                                                                                                                                                                                                                                                  Conservative 0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       (first entry)
                                                                                                                                                              (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      expressed sequence tag; secreted protein; cDNA isolation;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    99US-0122487P
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Duclert A,
                                                                                                                                                                                                                                                                                                                                                                                                     Score 52;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Giordano
                                                                                                                                                                                                                                                                                                                                                                                  1.8e-13;
hes 0;
                                                                                                                                                                                                                                                                                                                                                                                                                     DB 3;
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                                                                                                                                                                                                                                                                                                                                                                                                                     Length 381;
                                                                                                                                                                                                                                                                                                                                                                                    Indels
                                                                              ; forensic; disorder; ss.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                    Query Match
Best Local S
Matches 52
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess
                                                                                                                     Human; reproductive system related antigen; reproductive cancer; gene therapy; ds.
                                                                                                                                                                            Human reproductive system related antigen DNA SEQ ID NO: 8895.
                                                                                                                                                                                                                21-NOV-2001
                                                                                                                                                                                                                                                                                      AAL06207 standard; DNA; 9620 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 1437 BP;
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23-AUG-2000;
                                                                                     Homo sapiens.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                    Local Similarity nes 52; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                 3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                              243
                                                                                                                                                                                                                                                                                                                                                                              CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                    Conservative 0;
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                                                                                                                                                                                                                (first entry
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2000US-00649167.
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100.0%; Pred. No. 1.
ive 0; Mismatches
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                                                                                                                                          system disorder;
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02-AUG-2001 WO200155320-A2

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31-JAN-2000

24-FEB-2000)

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24-FEB-2000)

16-MAR-2000)

17-MAR-2000)

17-MAR-2000)

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2000US-0179065P.
2000US-018466P.
2000US-018466P.
2000US-018466P.
2000US-0198123P.
2000US-029467P.
2000US-0214866P.
2000US-0214866P.
2000US-021668P.
2000US-021668P.
2000US-021668P.
2000US-021668P.
2000US-021746P.
2000US-021746P.
2000US-0225214P.
2000US-0225214P.
2000US-0225214P.
2000US-0225214P.
2000US-0225266P.
2000US-0225270P.
2000US-022575P.
2000US-0231414P.
2000US-023144P.
2000US-023144P.
2000US-023144P.
2000US-023144P.
2000US-0231414P.
2000US-023149PP.
2000US-0231439P.
2000US-023364P.
2000U
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      29-SRP-2000
02-CCT-2000)
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03-CCT-2000)
03-CCT-2000)
03-CCT-2000)
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03-CCT-2000)
03-NOV-2000)
03-NOV
                                                                                            Rosen
    Isolated nucleic acid molecule encoding a reproductive system antigen
                                                                                                                                    (HUMA-)
                                               2001-465570,
                                                                                         ß
                                                                                         Barash
                                                                                                                                                                          2000US-0236370P.
2000US-023703PP.
2000US-0237039P.
2000US-0237039P.
2000US-0237039P.
2000US-0237049P.
2000US-0237049P.
2000US-0241966P.
2000US-0241808P.
2000US-0241808P.
2000US-0246471P.
2000US-0246471P.
2000US-0246477P.
2000US-0246477P.
2000US-0246477P.
2000US-0246478P.
2000US-0246523P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-024921P.
2000US-02511868P.
2000US-02511868P.
2000US-02511869P.
2000US-0251989P.
2000US-025199P.
2000US-025199P.
2000US-025499P.
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RESULT
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Best Local S
Matches 52
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                used
The present invention relates to methods for distinguishing between individuals homozygous for and therefore afflicted with Van Buchem's disease, individuals heterozygous for and therefore carriers of Van Buchem's disease and individuals who are not afflicted with Van Buchem's Buchem's disease comprise identifying a large genomic deletion in chromosome 17 at 17g21. The method is useful for identifying individuals who are afflicted with or carriers of diseases associated with one or more genomic
                                                                                                                                                                                                                                    Methods for identifying subjects who are afflicted with or carriers of diseases associated with genomic deletion(s), e.g. Van Buchem's disease by determining the presence of a deletion in the 92 kb region of human chromosome 17 at 17q21.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 07-FEB-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Homo
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human; Van Buchem's disease; genomic deletion; craniotubular hautosomal recessive disorder; chromosome 17; chromosome 17q21; bone dysplasia; 92Kb gene fragment; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Sequence
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                                                                                                                                                                                        Claim 14; Page 45-72; 109pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           28-JUL-2000; 2000US-0221855P
06-JUL-2001; 2001US-0303386P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  30-JUL-2001;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         AAD31364 standard;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                      (CELL-)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           the prevention and treatment of reproductive system disorders, cluding cancer. The present sequence is a genomic sequence enco
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                present invention provides the protein and
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                                                                                                                                                                                                                                                                                                                                                                                                                                                   CELLTECH R & D INC.
STRAEHLING HAMPTON K.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      9620
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             fragment in human chromosome 17
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Location/Qualifiers 5799. .57515
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            /*tag= a
/note= "This region is deleted in individuals afflicted
or carriers of Van Buchem's disease"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      2586 A; 2358 C;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ĕ
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                                                                                                                                                                                                                                                             Van Buchem's disease,
2 kb region of human
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           sequence encoding
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These can be used
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RESULT 22
ADF11613/G
ID ADF111
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Best Local S
Matches 52
                                                                               Determining a risk for or presence of altered bone mineral density (e. osteoporosis) in a subject comprises determining the presence or abser of a sclerostin gene region nucleotide polymorphism in a biological sample from a subject.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               deletion, particularly Van Buchem's disease, which is a rare recessive disorder that results in a bone dysplasia referred craniotubular hypertosis. The present sequence is a 92Kb gene
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           variation
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                                             Claim
                                                                                                                                                                                         WPI; 2003-833790/77.
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UNIV ROTTERDAM
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     92139 BP; 23017 A; 22243 C; 23264 G; 23612 T; 0 U; 3
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                                           ID NO 1;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  replace (74235,G)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           replace
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             replace (17966,G)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              replace(10565. .10566,AGGAC)
/*tag= c
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                                           114pp; English.
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        bone mineral density;
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                                                                                                                                                                                                                                  BW,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            dysplasia;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      gene fragment
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invention

relates to a method of determining a risk

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RESULT 23
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Best Local :
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 presence or absence of at least one sclerostin gene region nucleotide polymorphism in a biological sample from a subject where the presence of at least one polymorphism at a position that corresponds to a non-coding region of the 130320 bp sclerostin gene region (SOST) indicates an increased risk of altered BMD. The composition and methods are useful in determining in a subject a risk for having, or presence of, altered bone fracture or other conditions characterized by decreased or increased bone fracture or other conditions characterized by decreased or increased bone density. These may also be used in identifying agents that may be used for treating the above diseases, disorders or conditions associated with altered BMD. In addition, these may be used for pharmacogenomic purposes, e.g. to stratify patient populations according to suitability of a particular therapeutic agent for use in the population. This sequence corresponds to the human sclerostin gene region.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            SMRT inhibitor; cytostatic; antiinflammatory; antiarthritic; antirheumatic; antisense therapy; inflammatory disorder; rheumatoid arthritis; hyperproliferative disorder; cancer; leukaemia;
                                                                                                                                                                     Novel antisense compound targeted to nucleic acid encoding SMRT (silencing mediator for retinoid and thyroid hormone action), usefu treating animal having disease associated with SMRT such as cancer,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 SMRT; silencing mediator for retinoid and thyroid hormone action;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human SMRT partial genomic DNA sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Sequence 130320 BP; 33204 A; 32954 C; 31896 G; 32253 T; 0 U; 13 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       of altered bone mineral density (BMD) in a subject by determining
                                                                                                                                                                                                                                            GENBANK; NT_009459.
                                                                                                                                                                                                                                                                                                 Bennett CF,
                                                                                                                                                                                                                                                                                                                                                                    17-JUN-2002; 2002US-00174014.
                                                                                                                                                                                                                                                                                                                                                                                                     17-JUN-2003; 2003WO-US018923.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        11-MAR-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ADG86300
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ADG86300 standard; DNA; 220756 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  2889 GAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940
                                                                                                                                                                                                                                                                2004-082184/08
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               cancer;
                                                                                                                                                                                                                                                                                                                                    ISIS PHARM INC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Conservative
                                                                                                                                                                                                                                                                                                 Freier SM,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 human;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   100.0%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               gene; ds.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Score 52;
Pred. No.
                                                                                                                                                                                                                                                                                                 ₹
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     SEQ ID NO:14.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1.4e-13;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Length 130320;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Gaps
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targeted to a nucleic acid molecule encoding SMRT (silencing mediator for retinoid and thyroid hormone action), where (I) specifically hybridises with the nucleic acid molecule encoding SMRT and inhibits expression of SMRT. (I) specifically hybridises with at least 8-nucleobase portion of a preferred target region on nucleic acid molecule encoding SMRT. Also

present invention describes a compound (I) 8-50 nucleobases in length

Example 15;

SEQ ID NO 14; 260pp;

English.

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The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (1) for screening drug candidates; (11) for screening of bloactive agent capable of bioactive agent capable of bioactive agent capable of modulating the activity of CAP; (11) for screening of bloactive agent capable of modulating the activity of CAP; (1v) for carcinoma; (vi) for inhibiting the activity of CAP; (vi) for treating carcinoma; (vi) for neutralizing the effect of CAP; (vii) for treating C carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma hasociated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of Carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Matches 52;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Query Match
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              described is a composition (II) comprising (I) and a carrier or diluent.
(I) and (II) have cytostatic, antiinflammatory, antiarthritic and antirheumatic activities, and can be used in antisense therapy, and as SMRT expression inhibitors. (I) is useful for inhibitors that the expression of SMRT in cells or tissues. (I) is also useful for treating an animal having a disease or condition associated with SMRT, e.g., inflammatory disorder such as rheumatoid arthritis; or a hyperproliferative disorder such as cancer chosen from leukaemia and breast cancer, by inhibiting the expression of SMRT. (I) is useful for diagnostics, therapeutics, prophylaxis and as research reagents and kits. The present sequence represents a partial genomic DNA sequence of human SMRT, which is used in an example from the present invention. N.B. The present sequence is designated as SEQ ID NO:12 in example 15 but corresponds to SEQ ID NO:14
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Cytostatic; carcinoma; lymphoma; cancer; human; gene;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Human genomic sequence hCG25303.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ACN44282
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                                                                                                                                                                                                                                                                                                                    Claim 1; SEQ ID NO 652; Opp; English.
                                                                                                                                                                                                                                                                                                                                                          Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         28-FEB-2003; 2003WO-US006235
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                                                                                                                                                                                                                                                                                                                                                                                                                          2003-328604/31
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Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Matches 52;
                                                                                       isolation and characterisation of the DNA and protein sequences of the invention. The breast and ovarian cancer associated DNA, protein, agonist or antagonist sequences exhibit cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral; antiallergic; hepatotropic; antidiabetic; antinflammatory; antivicer; vulnerary; anticonvulsant; antibacterial; antifungal; antiparasitic and cardiant activity. The polynucleotide and protein sequences are used in the diagnosis of cancer, particularly breast and ovarian cancer. The nucleic acid sequences, proteins, agonists and agonists may also be used in the diagnosis, prevention and treatment of immune disorders e.g. Addison's disease, allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease, militing aclerosis proteins and ulcorative colities.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Human; breast cancer; ovarian cancer; cytostatic; immunosuppressive; nootropic; neurpprotective; antiviral; antiallergic; hepatotropic; antidiabetic; antiinflammatory; antiulcer; vulnerary; anticonvulsant; antibacterial; antifungal; antiparasitic; cardiant; immune disorder; Addison's disease; allergy; autoimmune haemolytic anaemia; autoimmune thyroiditis; diabetes mellitus; Crohn's disease; multiple sclerosis; rheumatoid arthritis; ulcerative colitis; cardiovascular disorder; wound healing; neurological disease; ds.
                                   disease, multiple sclerosis, rheumatoid arthritis and ulcerative colitis; cardiovascular disorders such as myocardial ischaemias; wound healing;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Sequences AAF21614 - AAF22031 represent DNA sequences encoding hupoteins AAB59111 - AAB59128. The DNA and protein sequences are associated with Dreast and ovarian cancer. Included in the invent sequences AAF22032 - AAF22040 and AAB59129 which are used in the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           New human breast and ovarian cancer associated gene sequences and the polypeptides encoded by these genes, useful in the prevention, treatmund and diagnosis of cancer, immune disorders, cardiovascular disorders as
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Rosen CA,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Ruben SM
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100.0%; Pr/
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0; Mismatches
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1.4e-13;
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                       cc sequences. (I) is useful as hybridisation probes, polymerase chain creation (PCR) primers, oligomers, and for chromosome and gene mapping, cc and in recombinant production of (II). The polymucleotides are also used civity of (II) or to treat disease states involving expressed cc genes. (I) is useful in gene therapy techniques to restore normal cc useful for generating antibodies against it, detecting or quantitating a cc polypeptide in tissue, as molecular weight markers and as a food csupplement. (II) and its binding partners are useful in medical imaging co f sites expressing (II). (I) and (II) are useful in medical imaging co sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The collegenties and polymucleotide sequences have applications in cc diagnostics, forensics, gene mapping, identification of mutations cc and to produce other types of data and products dependent on DNA and cambine active types of data and products dependent on DNA and cambine active types of the invention. Note: The sequence data for this collegent in the printed specification, but was obtained in celetronic format directly from MIPO at the print of the invention.
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Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                    New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess
                                                                                                                                                                                                                                                                                                                                                                                                  Claim 1; SEQ ID NO 29534; 103pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  31-MAR-2000;
23-AUG-2000;
            ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             P-PSDB;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             WPI; 2001-639362/73
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       30-MAR-2001; 2001WO-US008631.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 DNA encoding novel human diagnostic protein #29534.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     13-FEB-2002
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          infectious diseases
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2000US-00649167.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 INC.
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. 5.1e-13;
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    and polypeptide (II)

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                                                                                                                                                                                                                                                                                                                                                 The invention relates to a method (MI) for identifying one or more CC nucleic acid sequences useful as a biomarker for a disease to be detected. (MI) involves identifying nucleic acid sequences comprising CC detected. (MI) involves identifying nucleic acid sequences comprising CC regulated CpG site in promoter-first exon region and that are down-cc regulated in diseased cells, comparing expression level of nucleic acid sequences and controlled con
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Matches
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Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Identifying nucleic acid sequences as biomarker for disease, by identifying nucleic acid sequences comprising methylated CpG site and down regulated in diseased cells and comparing its expression level with
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Sequence 2791 BP; 559
                                                                                                                                                      nucleic acid sequences useful as a biomarker for a disease to be detected, where the nucleic acid sequences are useful for detecting, the presence or stage of a disease such as cancer e.g. colorectal cancer in a subject. The present sequence represents a specifically claimed human genomic sequence for use in the method of the invention. Note - The
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Beard C, Burgess C,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     16-DEC-2003; 2003US-00737082.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      (FARB ) BAYER CORP.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         DNA methylation; biomarker; cancer; gene; ds; SLC26A2.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        25-AUG-2005
                                                                                                                    sequence data
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          1538 CCTAAGGCGGAGCGCGCGCTCTGCAGCCTTGCCCCGGAGTTGGCACC 1588
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ANK; AI025519.
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                                                                               for this patent is not represented in the printed but was obtained in electronic format from the USI
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  1.6%;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Gannon A,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                0
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ВP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Harvey J,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               DB 5;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      4.9e-13;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Lechner JF,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Length 2791;
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Best Local Similarity
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                                                                                                                                                                                               14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   AAK67239;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 23139 BP; 6124 A; 4783 C; 4952 G; 7280 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             06-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                       31-JAN-2000;
04-FEB-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         AAK67239 standard; DNA; 30393 BP
                                                                                                                                                  14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                         14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                             07-JUN-2000;
28-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Human immune/haematopoietic antigen genomic sequence SEQ ID NO:22051.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              2890 AGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               51;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 AGGCAGGTGGATCACCTGAGGCCGAGGAGTTCGAGACCAGCCTGGCCAACAT 3733
2000US-0209467P.
2000US-0215135P.
2000US-0216647P.
2000US-0216880P.
2000US-021748FP.
2000US-021748FP.
2000US-022964P.
2000US-022964P.
2000US-0225213P.
2000US-0225214P.
2000US-0225214P.
2000US-0225268P.
2000US-0225268P.
2000US-0225447P.
2000US-0225447P.
2000US-0225479P.
2000US-0225479P.
2000US-0225759P.
2000US-0225759P.
2000US-022576681P.
2000US-022576681P.
2000US-0225768P.
2000US-0225768P.
2000US-0225768P.
2000US-0225768P.
2000US-0225759P.
2000US-02259349P.
2000US-0229344P.
2000US-0229344P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (first entry)
                                                                                                                                                                                                                                                                                                                                                                                    2000US-0190076P.
2000US-0198123P.
2000US-0205515P.
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Pred. No. 4.5e-13;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Length 23139;
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RESULT 29
ADX80722/c
ID ADX80722;
XX ADX80722;
AC ADX80722;
XX O5-MAY-200
XX DE Human manr
XX
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                                                                                                                                                                                                      S
                                                                                                                                                                                                                                                                                                     CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic contivity, and can be used in gene therapy and vaccine production. (I) cytoteins and polynucleotides may be used in the prevention, diagnosis and creatment of diseases associated with inappropriate (I) expression. For cerample, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome concert affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) concert affect caids into a host cell and culturing the cell to express the concert and treat immune/haematopoietic-related diseases, especially concerts and cancer metastases of haematopoietic acids expresent human immune/haematopoietic antigen genomic concerts and cancer metastases of haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK94950 and AAM82169 crepresent sequences used in the exemplification of the present invention.
                                                                                                                                                                                                                                      Query Match 1.6%;
Best Local Similarity 100.0%;
Matches 51; Conservative (
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    17-NOV-2000
17-NOV-2000
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17-NOV-2000;
17-NOV-2000;
              Human mannose receptor C type 2 (ENDO180)
                                              05-MAY-2005
                                                                                                                                                                                                                                                                                                   Sequence 30393 BP; 8271 A; 7315 C; 8076
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Disclosure; SEQ ID NO 22051; 3071pp + Sequence Listing; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Nucleic acids encoding useful for preventing,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                WPI; 2001-483426/52.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          (HUMA-)
                                                                                                                                                                              24742 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 24792
                                                                                                                                                                                                 3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 3121
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          HUMAN GENOME
                                                                                                     standard; DNA; 68200
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      2000US-0249218P
2000US-0249218P
2000US-0249244P
2000US-0249245P
2000US-0249264P
2000US-0249299P
2000US-0249299P
2000US-0249399P
2000US-0250160P
2000US-0251030P
2000US-0251030P
2000US-0251186P
2000US-0251869P
2000US-0251869P
2000US-0251869P
2000US-0251989P
2000US-0251989P
2000US-0251999P
2000US-0251999P
2000US-025199PP
2000US-025199PP
2000US-025199PP
2000US-025199PP
                                             (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   human immune/hematopoietic antigen diagnosing and/or treating cancers
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Ruben
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            INC.
                                                                                                                                                                                                                                     Score 51; DB; Pred. No. 4.5
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                                                                                                        BP
                                                                                                                                                                                                                                                      DB 4; Le
. 4.5e-13;
                                                                                                                                                                                                                                                                                                   G; 6731 T; 0 U; 0 Other;
               genomic
                                                                                                                                                                                                                                        0,
                                                                                                                                                                                                                                                                    Length 30393;
                                                                                                                                                                                                                                          Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     polypeptides, and metastasis.
                                                                                                                                                                                                                                          0;
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09-NOV-2000
01-NOV-2000

2000US-0231414P 2000US-023208BP 2000US-023208BP 2000US-0232339P 2000US-0232339P 2000US-0232340P 2000US-023240P 2000US-0233406P 2000US-0234223P 2000US-0234274P 2000US-0234274P 2000US-023499P 2000US-023499P 2000US-023499P 2000US-023499P 2000US-023499P 2000US-023499P 2000US-0235834P 2000US-0235834P 2000US-0235834P 2000US-0236368P 2000US-0236368P 2000US-024678P 2000US-0241868P 2000US-0241886P 2000US-0241878P 2000US-024677P 2000US-024677P 2000US-024677P 2000US-024677P 2000US-024652P 2000US-024652P 2000US-024652P 2000US-024652P 2000US-0246611P 2000US-0249211P 2000US-0249211P 2000US-0249211P 2000US-0249211P 2000US-0249211P 2000US-0249211P 2000US-0249211P

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ACN44754/c
ID ACN447
XX
AC ACN447
XX
DT 18-NOV
XX
Human
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Cytost
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OS Homo s
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                                                                                                                                                                                                                                                                                                                                                     S
                                                                                                                                                                                                                                                                                                                                                                                                                                                  of melanoma. The method comprises detecting the presence or absence of a copymorphic variation associated with melanoma, where the presence of the cone or more polymorphic variations is indicative of the subject being at risk of melanoma. The invention further comprises: a method for identifying a polymorphic variation associated with melanoma; an isolated nucleic acid which comprises a portion of or all of a nucleotide sequence comprising fully defined 68400-213300 base pairs sequences (SEQ ID NO. 3, 4, 5, 6, and/or 7) given in the specification, and comprises one or more polymorphic variations; an oligonucleotide comprising a nucleotide sequence complementary to a portion of the nucleotide sequence above, where the 3' end of the oligonucleotide comprising a nucleotide where the 3' end of the oligonucleotide comprising the isolated nucleic acid linked to a solid support; an isolated polypeptide encoded by the isolated nucleic acid sequence; genotyping a nucleic acid; a method for identifying a
                                                                                                                                                                                                                                                                                                                                                                                                           Query Match
Best Local
                                                                                                                                                                                                                                                                                                                                                                                        Matches 51;
 12-SEP-2003
                                   WO2003073826-A2
                                                                                                  Cytostatic; carcinoma; lymphoma; cancer; human;
                                                                                                                                   Human genomic sequence hCG37990.
                                                                                                                                                                      18-NOV-2004
                                                                                                                                                                                                       ACN44754;
                                                                                                                                                                                                                                     ACN44754 standard; DNA; 215221 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                The invention relates to a novel method for identifying a subject at
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Claim 16; SEQ ID NO 3; 418pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Identifying a subject at risk of melanoma by detecting presence or absence of a polymorphic variation associated with melanoma, where the presence of polymorphic variations is indicative of the subject being a risk of melanoma.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      WPI; 2005-182387/19.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      23-JUL-2003; 2003US-0489703P.
06-NOV-2003; 2003US-00703389.
06-NOV-2003; 2003US-00703817.
06-NOV-2003; 2003US-00704513.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            05-MAY-2004; 2004WO-US014238
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (SEQU-)
                                                                                                                                                                                                                                                                                                                                                                                                        Local Similarity
                                                                                                                                                                                                                                                                                                                       63277
                                                                                                                                                                                                                                                                                                                                       2890 AGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        SEQUENOM INC
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                                                                                                                                                                                                                                                                                                                       AGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 63227
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Nelson MR,
                                                                                                                                                                                                                                                                                                                                                                                    1.6%; Scilarity 100.0%; F
                                                                                                                                                                      (first entry)
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67995
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            standard_name= "Single nucleotide polymorphism"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Kammerer SM,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              a۷
                                                                                                                                                                                                                                                                                                                                                                                                        Score 51; DB 14;
Pred. No. 4.3e-13;
                                                                                                                                                                                                                                                                                                                                                                                        Mismatches
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                                                                                                  gene; ss
                                                                                                                                                                                                                                                                                                                                                                                                                      Length 68200;
                                                                                                                                                                                                                                                                                                                                                                                        Indels
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RESULT 31
AAC24464/c
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Best Local Similarity
Matches 51; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bioactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of bioactive agent capable of modulating the activity of CAP; (iv) for a bioactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing carcinoma; (vii) for inhibiting the activity of CAP; (vii) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent
                                                                                                                                                                                                                                                                            gene therapy;
                                                                                                                                                                                                                                                                                                                                                                                                            AAC24464 standard; cDNA; 255
                                             WPI; 2000-500381/45
                                                                                                                                 26-FEB-1999;
                                                                                                                                                             21-FEB-2000; 2000EP-00200610
                                                                                                                                                                                          06-SEP-2000
                                                                                                                                                                                                                       EP1033401-A2
                                                                                                                                                                                                                                                 Homo sapiens.
                                                                                                                                                                                                                                                                                           Human;
                                                                                                                                                                                                                                                                                                                        Human secreted protein 5' EST, SEQ ID NO: 28539.
                                                                                                                                                                                                                                                                                                                                                     06-OCT-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 215221 BP; 63216 A; 39385 C; 42715 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Claim 1; SEQ ID NO 1360; Opp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      01-MAR-2002; 2002US-00087192
                                                                        Dumas Milne Edwards
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     US2002182586A1,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             The present invention relates to novel DNA and protein sequences which
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  WPI; 2003-328604/31.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    28-FEB-2003;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (SAGR-) SAGRES DISCOVERY
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      50858 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 50808
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGACACTCTGTCT 3121
                                                                                                                                                                                                                                                                                           5' EST;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             1.6%; So ilarity 100.0%; I Conservative 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    2003WO-US006235
                                                                                                                                                                                                                                                                                                                                                     (first entry)
                                                                                                                                                                                                                                                                           chromosome mapping;
                                                                                                                                  99US-0122487P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  for which no sequence data was published
                                                                       'n
                                                                          Duclert
                                                                                                                                                                                                                                                                               mapping; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Score 51; DB; Pred. No. 4.2.
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                                                                        Giordano J;
                                                                                                                                                                                                                                                                                           secreted protein;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Length 215221;
tag (5' EST) for to 5'ESTs and for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Indels
                                                                                                                                                                                                                                                                                             cDNA isolation;
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New nucleic acid that is a 5' expressed sequence obtaining cDNAs and genomic DNAs that correspond

and for

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                cardiovascular; respiratory; gene therapy; secreted protein; chromosome identification; hybrid mapping; gene expression control; immune system disorder; immunodeficiency; Chediak-Hjgashi syndrome; autoimmune disease; systemic lupus erythematosus; rheumatoid arthritis; multiple sclerosis; haemolytic anaemia; myasthenia gravis; allergic reaction; asthma; inflammatory condition; inflammatory bowel disease; B cell stimulator; T cell activator; blood-related disorder; eosinophilia; thrombosis; thromboembolism; atherosclerosis; myocardial infarction; angina; anaemia; hyperproliferative disorder; cancer; renal disorder; chronic kidney failure; renal tubular acidosis, kidney stone;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               immunomodulator; immunosuppressive; antiinflammatory; dermatological; antiarthritic; antirheumatic; neuroprotective; antianaemic; muscular; antiallergic; antiasthmatic; gastrointestinal; anticosqulant; thrombolytic; antiarteriosclerotic; cardiant; cytostatic; nephrotropic;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  diagnostic, forensic, gene therapy and chromosome mapping procedures.
                                                                                                                                                                                                                                                                              10-OCT-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             cardiovascular
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Novel human
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08-JUL-1997;
08-JUL-1997;
08-JUL-1997;
                     anemia,
kidney f
                                                                                                  Fischer
Lafleur
                                New isolated nucleic acid encoding human preventing or diagnosing e.g. rheumatoid anemia, inflammatory bowel disease, ather
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ROSEN C A.
SOPPET D R.
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OLSEN H.
EBNER R.
BIRSE C F
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SEQ ID
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Moore
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Shi Y, Olsen
                                                                                                  Ruben SM, Kyaw H, Li
sen H, Ebner R, Birse
                                   atherosclerosis, cancers,
                                           proteins, useful for treating, arthritis, multiple sclerosis,
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                                                                                                 GE,
γ,
                                   chronic
                                                                                                              Zeng
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The invention describes novel human secreted proteins and the nucleotides encoding them. The polynucleotides are useful in chromosome identification, for radiation hybrid mapping, in controlling gene expression, in gene therapy or as molecular weight markers. The polynucleotides and polypeptides are useful for diagnosing, treating or preventing diseases of the immune system, immunodeficiencies, e.g. Chediak-Higashi syndrome, autoimmune diseases, e.g. systemic lupus erythematosus, rheumatoid arthritis, multiple sclerosis, haemolytic anaemia or myasthenia gravis, allergic reactions, e.g. asthma, inflammatory conditions, e.g. inflammatory bowel disease. They can also be used as a stimulator of B cell responsiveness to pathogens or as an activator of T cells. The polynucleotides and polypeptides are also useful for treating or preventing blood-related disorders, e.g. eosinophilia, thrombosis, thromboembolism, atherosclerosis, myocardial infarction, unstable angina or anaemia. They can also be used for treating, preventing or diagnosing hyperproliferative disorders (chronic kidney failure, renal tubular acidosis or kidney stones), cardiovascular disorders or respiratory disorders. This sequence represents a novel human secreted protein

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RESULT 33
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24-FEB-2000;
16-MAR-2000;
11-MAR-2000;
11-MAR-2000;
11-MAR-2000;
28-JUN-2000;
28-JUN-2000;
20-JUL-2000;
11-JUL-2000;
11-AUG-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              polynucleotide fragment. Note: This sequence is available format from the US patent office at ftp.seqdata.uspto.gov/sequence.html?DocID=20040044191.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   17-JAN-2001; 2001WO-US001354
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nes 50; Conservative (
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2000US-0179065P
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2000US-0214866P
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2000US-022575PP
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2000US-023239P

2000US-023423P

2000US-0235836P

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2000US-024651P

2000US-024651P

2000US-024651P

2000US-024621P

2000US-024651P

2000US-024651P
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17-NOV-2000; 17-NOV-2000; 17-NOV-2000; 17-NOV-2000; 17-NOV-2000;

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RESULT 34
AAS93725
ID AAS93
XX
AC AAS93
XX
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                                                                                                                                                                                                 Query Match
Best Local
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17-NOV-2000
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05-DEC-2000
05-DEC-2000
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06-DEC-2000
08-DEC-2000
08-DEC-2000
                                                                                                                                                                                                                                                                    example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, diagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metastases of haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoietic antigen genomic sequences from the present invention. AAK84942 to AAK84950 and AAM82169 represent sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                      amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            17-NOV-2000;
17-NOV-2000;
               AAS93725;
                                                                                                                                                                                                                                           Sequence 301 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 38904; 3071pp + Sequence Listing; English.
                                             AAS93725 standard; cDNA; 432 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              WPI; 2001-483426/52.
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                                                                                                                        234 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                 Similarity
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2000US-0251030P.
2000US-025198BP.
2000US-0256719P.
2000US-0251879P.
2000US-0251868P.
                                                                                                                                                                                 Conservative
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2000US-0254097P
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2000US-0251989P.
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2000US-0250160P.
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2000US-0249299P.
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                                                                                                                                                                                                                                           A; 74 C; 92
                                                                                                                                                                                                               1.6%;
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                                                                                                                                                                                 0
                                                                                                                                                                                              Score 50;
Pred. No.
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                                                                                                                                                                                 Mismatches
                                                                                                                                                                                                                                           48 T; 0 U;
                                                                                                                                                                               1.5e-12;
hes 0;
                                                                                                                                                                                                             DB 4;
                                                                                                                                                                                                                                             0 Other;
                                                                                                                                                                                                            Length 301;
                                                                                                                                                                                 Indels
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04-NOV-2004 ADQ81170; ADQ81170

(first entry)

standard;

DNA; 1001

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RESULT 35
ADQ81170
ID ADQ81
XX
AC ADQ81
XX
DT 04-NC
XX
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ci in diagnostics as expressed sequence tags for identifying expressed couseful for generating antibodies against it, detecting or parantitating a colvity of (II) or to treat disease states involving (II). (II) is cuseful for generating antibodies against it, detecting or quantitating a colving country of the protein expression of the sequence and as a food complement. (II) and its binding partners are useful in medical imaging cof sites expressing (II). (I) and (II) are useful for treating disorders covolving aberrant protein expression or biological activity. The complement of the invention of the triatist of assess biodiversity and to produce other types of data and products dependent on DNA and compared to produce other types of data and products dependent on DNA and coming sequences of the invention. Note: The sequence data for this catenit did not appear in the printed specification, but was obtained in clectronic format directly from WIPO at control of the country pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                   Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Query Match
Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess biodiversity.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 432 BP; 57 A; 142 C; 168 G; 65 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polymolectides are also used
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               WPI; 2001-639362/73.
P-PSDB; ABG29538.
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23-AUG-2000; 2000US-00649167.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       DNA encoding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Claim 1; SEQ ID NO 29529; 103pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Drmanac RT,
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                                                                                                                                                                                                                                                                                                                                              888 GACGGGCTCAACAAGACGACTGCGTGCTACCACCACCTGGTGCTGACCGT
                                                                                                                                                                                                                                                                                                               22
                                                                                                                                                                                                                                                                                                                                                                                                                                   50;
                                                                                                                                                                                                                                                                                                               GACGGGCTCAACAAGACGACTGCGTGCTACCACCACCTGGTGCTGACCGT
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                                                                                                                                                                                                                                                                                                                                                                                                                                Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                   0,
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                       Mismatches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Length 432;
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                                                                                                                                                                                                                                                                                                                                                                              937
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The invention relates to a novel polymucleotide encoded by a phenotype CC associated (PA) gene. The polymucleotide is selected from 292 sequences CC comprising 301-1002 base pairs (AD081204) given in the CC specification, with allelic variation contained in a functional CC surrounding like full length cDNA for PA gene polypeptide and with or CC without the PA gene promoter sequence. A polymucleotide of the invention has cardiant activity, and acts as a phenotype-associated gene modulator. The reagent of the invention is useful for preparing a medicament The CC method of the invention is useful for preparing a medicament tailored to suit a patient's individual response to statin therapy. The genetic CC polymorphisms are useful for assessing the response to lipid lowering CC drug therapy and adverse drug reactions of the medicaments, particularly for assessing cardiovascular risks in humans e.g. atherosclerosis, cischaemia/reperfusion, and stroke. The genetic polymorphisms are also useful for identifying compounds for treatments of cardiovascular disease compounds for treatments of cardiovascular disease and as templates for the recombinant production of normal variant peptides or polypeptides of the recombinant production of normal variant peptides or polypeptides of the genes. The present sequence represents a polynucleotide of the present sequence.
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                                    S
                                                                                 Query Match
Best Local S
Matches 50
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             New polymorphisms of a phenotype associated (PA) gene, useful for assessing the response to lipid lowering drug therapy and adverse drug reactions of the medicaments, and for screening compounds for treating
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          WPI; 2004-581012/56.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     31-JAN-2003; 2003EP-00002212.
03-FEB-2003; 2003EP-00002153.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      atherosclerosis; ischaemia; reperfusion; hypertension; restenosis; arterial inflammation; myocardial infarction; stroke; single nucleotide polymorphism; SNP.
                                                                                                                                                                     Sequence 1001 BP; 363
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      23-JAN-2004; 2004WO-EP000539
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human phenotype associated polynucleotide baySNP59113
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                variation
                                    3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                sapiens.
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                                                                                                     Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  SEQ ID NO 258; 349pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            phenotype associated; PA; cardiant; statin;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Schwers S,
                                                                                 Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Location/Qualifiers
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/standard_name= "Single nucleotide polymorphism"
                                                                          1.6%; but
100.0%; Pr
                                                                                                                                                                     A; 150 C;
                                                                               Score 50; DB;; Pred. No. 1.5:
                                                                                                                                                                     269 G;
                                                                                                                                                                     218 T; 0 U; 1 Other;
                                                                                 1.5e-12;
hes 0;
                                                                                                                            DB 13;
                                                                                                                          Length 1001;
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                                                                                 Gaps
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RESULT 36

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RESULT 37
ABA82624
ID ABA82
XX
AC ABA82
XZ
AC ABA82
XX
DT 25-JF
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Humar
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Best Local Similarity
Matches 50; Conserva
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Mao
                                                                                                                                                                                                                                                                                                                                     described is the process for preparing the protein by DNA recombination and the application of the polypeptide and polymucleotide in treating various diseases such as malignant tunours, haemopathy, human immunodeficiency virus (HIV) infection, immunological diseases, and various inflammations. The present sequence encodes human zinc finger protein 10.01
Human; high bone mass; HBM gene; Zmax1 gene; chromosome 11; 11q13.3; sequence tagged site; STS; osteoporosis; osteopathic; gene therapy;
                                        Human HBM gene region b200e21-h_contig1.
                                                                                                                                                                                                                                                                                                         Sequence 2407
                                                                                                                                                                                                                                                                                                                                                                                                                        The present invention relates to the isolation of human zinc finger protein 10.01, and the polynucleotide sequence encoding it. Also
                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 6; Page 25-26 (disclosure); 33pp; Chinese.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              diseases
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; zinc finger protein 10.01; malignant tumour; haemopathy;
human immunodeficiency virus infection; HIV infection; inflammation;
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                                                                                                                            ABA82624 standard; DNA; 8705 BP
                                                                                                                                                                                                                                                                                                                                                                                                                       protein 10.01,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            New human zinc finger protein 10.01 polypeptide tumors, hemopathy, human immunodeficiency virus
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     CN1352110-A.
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DB; ABG72222.
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                                                                                                                                                                                                                           AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              and various inflammations.
                                                                                                                                                                                                                                                   Conservative
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                                                                    (first entry)
                                                                                                                                                                                                                                                                                                           BP;
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1279. .1554
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                                                                                                                                                                                                                                                                                                           746 A; 458 C; 570 G; 633 T; 0 U; 0 Other;
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100.0%; Pred. No.
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                                                                                                                                                                                                                                                                              Length 2407
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            for treating malignant infection, immunological
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WWXSXEX EXEX EXX EXX EXX SOCOOO
ID XXX PR PR XX PR XX PR XXX PR PR PR XX P
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Best Local S
Matches 50
                      11-MAY-2001;
17-MAY-2001;
01-FEB-2002;
04-MAR-2002;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                The present invention describes the human Zmax1 gene and the high bone mass (HBM) gene, which are found on chromosome 11q.13.3. The Zmax1 and HBM genes have osteopathic activities. The genes can be used in gene therapy, antisense therapy and in the production of vaccines. They can be used in the diagnosis and treatment of bone disorders including osteoporosis, paget's disease, sclerostosis, osteomaliacia and fibrous dysplassia.

ABA82038 to ABA82700 and AAG68168 to AAG68193 represent sequences used in
                                                                                                                                                                                                                                                                                                                                                                                Human; high bone mass; HBM; LRP5; LRP6; transgenic; bone mass modulation; gene therapy; bone density modulation; bone strength; trabecular number; bone size, bone tissue connectivity; bone disease; osteoporosis; osteoparaia; rickets; Paget's disease; neoplasm of the bone; gene; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            05-APR-2000; 2000US-00543771.
05-APR-2000; 2000US-00544398.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   antisense therapy; vaccine; bone disorder; Paget's disease; sclerostosis; osteomalacia; fibrous dysplasia; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Carulli
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               21-JUN-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Homo
                                                                                                                                                               13-MAY-2002;
                                                                                                                                                                                                                                                                         WO200292764-A2
                                                                                                                                                                                                                                                                                                                                  Homo
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Human HBM
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               02-JUN-2003
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ACC45365
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            ACC45365 standard; DNA; 8705 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence 8705 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Mew high bone mass (HBM)
modulating bone mass for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (GENO-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  6492
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                             sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         l Similarity
50; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            GENOME THERAPEUTICS CORP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  ### AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCCAAGAGCTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      gene fragment #6.
                         ; 2001US-0290071P.
; 2001US-0291311P.
; 2002US-0353058P.
; 2002US-0361293P.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Little
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  2000WO-US016951.
                                                                                                                                                               2002WO-US014876.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      303-308; 443pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   2107
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   100.0%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         1.6%; Score 50; DB
100.0%; Pred. No. 1.
ive 0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   A; 2317 C; 2399 G; 1882
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          and Zmax1 genes and proteins useful the treatment of e.g. osteoporosis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Recker RR,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Johnson
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         1.4e-12;
hes 0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   T; 0
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      3122
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RESULT 39
ADB98065
ID ADB98
XX ADB98
XX ADB98
XX ADB98
XX OA-DE
XX OStec
KW Oone
XX Oone
XX Homo
OS Homo
XX Y 11-W2
PR 11-W2
PR 11-W2
PR 01-FE
PR 04-W3
XX OGENC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          The invention relates to novel transgenic animals expressing the high CC bone mass (HBM) gene, expressing the corresponding wild type HBM gene, comprising an alteration of the gene encoding LRP5 or LRP6, or expressing CC an LRP5 that is modulated by an altered gene control sequence introduced by homologous or non-homologous recombination. The transgenic animals are CC for the study of bone density modulation or bone mass modulation. The CC invention has osteopathic and cytostatic activity. The polynuclectides of the invention may have a use in gene therapy. The transgenic animals and CC uncleic acids are for the study of bone density modulation, where the CC bone mass is modulated relative to non-transgenic animals of the same CC strength, trabecular number, bone size, or bone tissue connectivity. The CC transgenic animals, nucleic acids and methods are useful for identifying molecules involved in bone development, and for developing pharmaceutical compositions, which may be employed for treating or preventing bone CC diseases, e.g. osteoporosis, osteomalacia, rickets, Paget's disease, or neoplasms of the bone. The transgenic animals and nucleic acids are also useful in methods for diagnosing diseases involved in bone development, or characterised by reduced bone density or mass. The present sequence is
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Query Match
Best Local 9
                                      11-MAY-2001; 2001US-0290071P.
17-MAY-2001; 2001US-0291311P.
01-FEB-2002; 2002US-0353058P.
04-MAR-2002; 2002US-0361293P.
                                                                                                                                                                                                                                                                                           Osteopathic; Gene therapy; High Bone Mass; HBM; LRP5; Zmax1; LRP6; bone mass modulation; osteoporosis; human; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 8705 BP; 2107 A; 2317 C; 2399 G; 1882 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             New transgenic animals (e.g. mice), useful as models for studying bone density modulation, developing drugs for treating or preventing bone diseases (e.g. osteoporosis), or diagnosing diseases characterized by
                                                                                                                                                                                                                                                                                                                                                     HBM-related clone contig b200e21-h contig1.
                                                                                                                                                                                                                                                                                                                                                                                           04-DEC-2003
                                                                                                                                                                                                                                                                                                                                                                                                                                                                       ADB98065
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Example 2; Page 358-361;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                reduced bone density.
                                                                                                                                      13-MAY-2002; 2002WO-US014877
                                                                                                                                                                              21-NOV-2002
                                                                                                                                                                                                                  WO200292000-A2.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       (GENO-) GENOME THERAPEUTICS (AMHP) WYETH.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                     standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              1.6%; Sc
llarity 100.0%; P
Conservative 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                         DNA;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        603pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Score 50; Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Bodine PV;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      DB 8;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Length 8705;
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(GENO-)

GENOME THERAPEUTICS CORP.

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RESULT 40
ADE82434
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Best Local S
Matches 50
                                                                                                                  Allen K,
Yaworsky
                                          Regulating LRP5, LRP6 or HBM activity in a subject, useful for modulating lipid levels and/or bone mass, and for in treating bone mass disorders, e.g. osteoporosis, comprises administering a composition which modulates
                                                                                                                            Allen
                                                                                                                                                                                     17-MAY-2001; 2001US-0291311P.
01-FEB-2002; 2002US-0353058P.
04-MAR-2002; 2002US-0361293P.
                                                                                                                                                                                                                                      17-MAY-2002; 2002WO-US015982.
                                                                                                                                                                                                                                                               21-NOV-2002.
                                                                                                                                                                                                                                                                                      WO200292015-A2
                                                                                                                                                                                                                                                                                                             Homo sapiens.
                                                                                                                                                                                                                                                                                                                                      hyperostosis;
                                                                                                                                                                                                                                                                                                                                                                                    Human DNA sequence
                                                                                                                                                                                                                                                                                                                                                                                                            29-JAN-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                   ADE82434;
                                                                                                                                                                                                                                                                                                                                                                                                                                                        ADE82434 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 8705
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       phenotype in a subject and for preparing a composition for modulating bone mass and/or lipid levels in a subject suffering from e.g. osteoporosis. The present sequence was used to illustrate the invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Example 3; SEQ ID NO 10; 629pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             suffering
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New nucleic acid comprising a mutation in LRP5 or LRP6, useful diagnosing a HBM-like phenotype in a subject and for preparing composition for modulating bone mass and/or lipid levels in a f
                                                                                          WPI; 2003-129219/12
                                                                                                                                                                                                                                                                                                                                                 Antiarthritic;
                                                                                                                                                                                                                                                                                                                                                            LRP5; LRP6; HBM; Dkk activity; Osteopathic; Antiinflammatory;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         (AMHP ) WYETH.
                                                                                                                                                                (GENO-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             evel modulation.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        present invention relates to High Bone Mass (HBM), LRP5 (Zmax1) and present invention relates to High Bone Mass (HBM), LRP5 (Zmax1) and 5 mutants, which results in a HBM-like phenotype when expressed in a 1. The HBM-like phenotype results in bone mass modulation and/or lipid rel modulation. The invention is useful for diagnosing a HBM-like rel modulation. The invention is useful for diagnosing a HBM-like
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     6492
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            3073
                                 osteoporosis, comprises k activity.
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                                                                                                                                                    GENOME '
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Similarity
                                                                                                                  Anisowicz PJ;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             from e.g. osteoporosis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Anisowicz
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Conservative
                                                                                                                                                                                                                                                                                                                                                                                                          (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  B₽;
                                                                                                                                                                                                                                                                                                                                     osteogenesis;
                                                                                                                                                                THERAPEUTICS
                                                                                                                                                                                                                                                                                                                                                 bone mass disorders;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               2107 A; 2317 C;
                                                                                                                                                                                                                                                                                                                                                                                    related
                                                                                                                                                                                                                                                                                                                                                                                                                                                          DNA;
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                                                                                                                                                                                                                                                                                                                                     signaling;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Mismatches
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                                                                                                                             Damagnez
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                                                                                                                                                                                                                                                                                                                                      osteoporosis; hypercalcaemia;
naling; ds.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  G; 1882
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             1.4e-12;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       DB 10;
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                                                                                                                             Robinson
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Length 8705;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Yaworsky
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Disclosure; SEQ

IJ

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10;

173pp; English

The invention relates to an isolated amino acid protein sequence selected from an amino acid sequence appearing as ADR16922 or an amino acid sequence comprising or consisting of the extracellular domain of ADR16922 (amino acids 23-1385). ADR16922 is encoded by the HBM (high bone mass) allele of the human Zmax1 gene and has sequence similarity to LDL receptors. Also disclosed are nucleic acids, proteins, cloning vectors, expression vectors, transformed hosts, methods of developing

Example 2; SEQ ID NO 10; 284pp; English.

pharmaceutical compositions, methods of identifying molecules involved in bone development, and methods of diagnosing and treating diseases involved in bone development. Specifically disclosed is the Zmax1 gene

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Query Match
                                                                                                                                                                                                                                                                                                                            13-JAN-1998;
23-OCT-1998;
13-JAN-1999;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        The present invention relates to regulating LRP5, LRP6 or HEM activity is a subject comprising administering a composition which modulates a Dkk activity. The method is useful for modulating lipid levels and/or bone mass, and is useful in treating or diagnosing abnormal lipid levels and bone mass disorders, such as osteoporosis, bond fracture, age-related loss of bone, a chondrodystrophy, drug-induced bone disorder, high bone turnover, hypercalcaemia, hyperostosis, osteogenesis, imperfects, osteomalacia, osteomyelitis, Paget's disease, osteoarthritis, and rickets. Modulators of Dkk activity are useful for as reagents in studying bone mass and lipid level modulation, in modulating Wnt signaling, or treating Dkk-mediated disorders. The present sequence represents a human DNA sequence related to the invention.
                                                                                                                                                                                     New high bone mass gene of chromosome 1.1Q13.3, encoding protein useful for treating, diagnosing, preventing, or screening for normal and abnormal conditions of bone, including metabolic bone diseases, e.g.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human; high bone mass; Zmax1; ds; BAC; HBM; osteoporosis; chromosome 11q13.3; osteopathic; LDL receptor; bone devel metabolic bone disease; bacterial artificial chromosome.
                                                                                                                                                                                                                                              WPI; 2004-623529/60.
                                                                                                                                                                                                                                                                      Carulli JP,
                                                                                                                                                                                                                                                                                                                                                                                   05-APR-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                BAC clone containing
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                                                                                                                                                                                                                                                                                                  (GENO-) GENOME THERAPEUTICS CORP.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   standard; DNA; 8705
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                                                                                                                                                                                                                                                                                                                                                                                   2000US-00543771.
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98US-0105511P.
99US-00229319.
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               high bone mass protein (HBM). The gene exists in two alleles, Zmaxi, the contional wild-type (the cDNA for which appears as ADR47570 encoding ADR47572) and the HBM allele (the cDNA for which appears as ADR47571 encoding ADR47573). The two alleles differ by a single nucleotide polymorphism (G to T at position 582 of ADR47570) causing a Gly to Val change at position 171 of the protein. Also included are a replicative claning vector comprising HBM/Zmaxi (and a replican operative in an isolated host cell), an expression vector comprising HBM/Zmaxi operably linked to a transcription regulatory region, an isolated host cell transformed with the vector(s), a method for testing a substance as a therapeutic agent for bone modulation in a host, a method of identifying a molecule involved in bone modulation, a method for testing for HBM activity, a method of developing a pharmaceutical for the treatment of bone development disorders, a method for treating a bone development in a host.
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Best Local S
Matches 50
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Sequence 8705 BP; 2107 A; 2317 C; 2399
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05-APR-2000; 2000US-00544398.
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23-OCT-1998;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Carulli JP,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           10-DEC-2003;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            bone modulation; bone development disorder; osteoporosis; chromosome 11q13.3; gene therapy; BAC.
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      in an
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          and/or preventing osteoporosis
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   SEQ ID NO 10; 303pp; English.
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98US-0105511P.
      animal,
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a method of
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Pred. No.
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      altering bone development
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Johnson
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           bone
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           mass,
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      in a host,
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RESULT 43
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  13-JAN-1998;
23-OCT-1998;
13-JAN-1999;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         CREIGHTON SCHOOL MEDICINE
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   bone
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        98US-0105511P.
99US-00229319.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                98US-0071449P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             injury; Pagets disease; ds
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1.6%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Recker RR, Johnson ML;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         °,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Score 50;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      1.4e-12;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Length 8705;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               6541
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The present invention relates to a method (M1) for identifying a candidate molecule involved in bone modulation. The method comprises identifying a molecule that binds to High Bone Mass protein (HBM) and/Zmax1 protein. The HBM gene exists in two alleles: Zmax1, the notional wild-type (the cDNA for which appears as AEB6929 encoding AEB69301 an AEB69300 encoding AEB69302). The two alleles differ by a single nucleotide polymorphism (T to G at position 582 of AEB69299)

and ď Example

2; SEQ ID NO 10; 308pp;

English.

Identifying candidate molecule involved in bone identifying molecule that binds to Zmax1, high because the control of the cont

high bone mass (HBM)

protein,

modulation,

Zmax1 and HBM

protein.

causing a Gly

6

change at position 171

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RESULT 44
AAK86119
ID AAK866
XX AAK86
XX AAK86
XX AAK86
XX Human
XX Human
XX Human
XX Human
XX Homo
XX
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  31-JAN-2000;
04-FBB-2000;
24-FBB-2000;
02-MAR-2000;
11-MAR-2000;
11-MAR-2000;
11-MAR-2000;
07-JUN-2000;
07-JUL-2000;
07-JU
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      protein has the property of causing elevated bone mass, while the Zmax1 protein does not. The gene for HBM/Zmax1 is located on chromosome 11q13.3. Also claimed is a method of pharmaceutical development for treating of bone development disorders, such as osteoporosis, osteomalacia, bone fractures, Paget's disease, etc., which comprises identifying a molecule that binds to the Zmax1 protein, or to HBM, or both. The present sequence was used to illustrate the invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         09-AUG-2001.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       cytostatic;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Human; immune; haematopoietic; immune/haematopoietic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Human immune/haematopoietic antigen
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         07-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                AAK86119;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Sequence 8705 BP; 2107 A; 2317 C; 2399
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              50;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    standard;
2000US-0179065P.
2000US-0184664P.
2000US-0184664P.
2000US-018465P.
2000US-0199076P.
2000US-0199076P.
2000US-0199076P.
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2000US-02252647P.
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2000US-0225758P.
2000US-0225758P.
2000US-0225758P.
2000US-0225758P.
2000US-0225758P.
2000US-0225768P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        1.6%; Score 50; DB ilarity 100.0%; Pred. No. 1.. Conservative 0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   2001WO-US001354.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       (first
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     therapy; vaccine; metastasis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    10396
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               genomic sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  G; 1882 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            DB 14;
1.4e-12
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          .4e-12;
s 0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 SEQ ID NO:40931.
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      22-AUG-2000
23-AUG-2000
30-AUG-2000
01-SEP-2000
01-SEP-2000
06-SEP-2000
06-SEP-2000
06-SEP-2000
08-SEP-2000
08-SEP-2000
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2000US-0246611P.
2000US-024652P.
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                                   Ś
                                                                                                                                                                                                              CC anino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic antiyity, and can be used in gene therapy and vaccine production. (I) coroteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For cc example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome cc that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) cc polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the corotein. (I) proteins and polynucleotides may be used to produce the secreted (I), by inserting the corotein. (I) proteins and polynucleotides may be used to prevent, cc diagnose and treat immune/haematopoietic-related diseases, especially cc cancers and cancer metastases of haematopoietic antigen genomic to AAX87694 represent human immune/haematopoietic antigen genomic the corotein invention. AAX54942 to AAX54950 and AAM82169
                                                                               Matches
                                                                                                Query Match
Best Local
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01-DEC-2000;
01-DEC-2000;
05-DEC-2000;
05-DEC-2000;
06-DEC-2000;
08-DEC-2000;
                                                                                                                                                            Sequence 10396
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 40931; 3071pp + Sequence Listing; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Nucleic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Rogen
  5597
                                       3073
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                                                                                                  Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         acids encoding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     HUMAN
AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 5646
                      AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                  sequences used in the exemplification
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         preventing,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Barash
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2000US-025198P.
2000US-0256719P.
2000US-0251479P.
2000US-0251856P.
2000US-0251869P.
2000US-0251869P.
2000US-0251989P.
2000US-0251999P.
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2000US-0249213P
2000US-0249215P
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2000US-0249299P
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2000US-0249299P
2000US-0249299P
2000US-0250160P
                                                                               Conservative
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2000US-0249210P.
2000US-0249211P.
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                                                                                                                                                          BP; 3175 A;
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                                                                                                1.6%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       SCI
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       human immune/hematopoietic antigen polypeptides, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Ruben SM;
                                                                                                                                                            2158 C;
                                                                               0;
                                                                                                Score 50; pred. No.
                                                                               Mismatches
                                                                                                                                                            2127
                                                                                                                       協
                                                                                                                                                          G; 2936 T; 0 U; 0 Other;
                                                                                                1.4e-12;
                                                                                                                       4.
                                                                               0
                                                                                                                  Length 10396;
                                                                                                                                                                                                  of the present invention
                                                                               Indels
                                                                           0;
                                                                           Gaps
                                                                             0
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Human nervous system related polynucleotide SEQ ID
                                                                                                        23-JAN-2002
                                                                                                              ABA20857 standard; DNA; 11234
                                                                                                        (first entry)
                                                                                                              BP.
                                                                                                     NO 13188.
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Human; nootropic; neuroprotective; cytostatic; dermatological; virus immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnes antiparkinsonian; antisickling; antianaemic; antiarthritic; cancer; antitheumatic; hepatorropic; cerebroprotective; antiinflammatory; antiallergic; antidiabetic; antiulcer; anticonvulsant; antifungal; antiparasitic; cardiant; immune disorder; cardiovascular disorder; antiparasitic; cardiant; immune disorder; cardiovascular disorder; neurological disease; infection; nephrotropic; gene therapy; vaccine; dermatological; virucide; antibacterial; vulnerary; ds.

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28-JUN-2000

30-JUN-2000

07-JUL-2000

07-JUL-2000

11-JUL-2000

14-JUL-2000

14-JUL-2000

14-JUL-2000

14-JUL-2000

14-JUG-2000

15-JUG-2000

16-JUG-2000

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18-JUG-2000

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17-MAR-2000;
18-APR-2000;
19-MAY-2000;
07-JUN-2000;
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24-FEB-2000;
02-MAR-2000;
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2000US-0184664P

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RESULT 45 ABA20857

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20-OCT
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08-SEP-2000;
12-SEP-2000;
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2000US-024929P
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2000US-0231968P.
2000US-023196P.
2000US-0232397P.
2000US-0232399P.
2000US-0232399P.
2000US-0232399P.
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                                                                  Query Match 1.6%; So
Best Local Similarity 100.0%; F
Matches 50; Conservative 0;
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01-DEC-2000
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05-DEC-2000
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08-DEC-2000
08-DEC-2000
08-DEC-2000
                                                                 Sequence 11234 BP; 3094 A; 2417 C; 2869 G; 2854 T; 0 U; 0 Other;
                                                                                                                                                                                                                                      Disclosure; SEQ ID NO 13188; 1701pp +
                                                                                  from WIPO at
                                                                                                                                                                                                                                                                useful for
                                                                                                                                                                                                                                                                        Nucleic acids encoding
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11155
                                                                                                                                                                                                                                                      rul for preventing,
metastases.
               3073
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        AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGACAGAGACTCTGTCTC 3122
AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 11204
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2000US-0259331P.
2000US-0251130P.
2000US-0251988P.
2000US-025189P.
2000US-0251858P.
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2000US-0251858P.
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2000US-0251869P.
2000US-0251869P.
2000US-0251869P.
2000US-0251869P.
                                                                                 ftp.wipo.int/pub/published_pct_sequences
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                               Score 50; DB:; Pred. No. 1.4:
0; Mismatches
                                                                                                                                                                                                                                      Sequence Listing;
                                        DB 5; Le
                                 <u>,</u>
                                                Length 11234;
                                 Indels
                                                                                                                                                                                                                                       English
                                <u>,</u>
                                Gaps
                                                                                                                                                                                                                                                                cancers
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RESULT 46
AAK80184/c
Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                        07-NOV-2001
                                                                                                                              AAK80184;
                                                                                                                                                 AAK80184
         WO200157182-A2
                            Homo sapiens.
                                                                                  Human immune/haematopoietic antigen genomic sequence SEQ ID NO:34996
                                                                                                                                                standard; DNA; 13026
                                                                                                       (first entry)
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SEP	26-SEP-2000;	21-SEP-2000; 25-SEP-2000; 25-SEP-2000;	14-SEP-2000; 14-SEP-2000; 21-SEP-2000;	14-SEP-2000;	14-SEP-2000;	14-SEP-2000;	08-SEP-2000;	08-SEP-2000;	08-SEP-2000;	08-SEP-2000;	06-SEP-2000;	05-SEP-2000;	01-SEP-2000;	01-SEP-2000;	01-SEP-2000;	30-AUG-2000;	22-AUG-2000;	22-AUG-2000; 22-AUG-2000;	18-AUG-2000;	4-AUG	4-AUG	14-AUG-2000; 14-AUG-2000;	4-AUG	4-AUG	14-AUG-2000; 14-AUG-2000;	4-AUG-200		1-JUL-2	11-JUL-2000;	7-JUL-2000	30-JUN-2000;	7-JUN-2	-MAY-	7-MAR-2		4-FEB-200	JAN-2	17-JAN-2001;	09-AUG-2001.
2000US-0236327P. 2000US-0236367P. 2000US-0236367P.	2000US-0235484P. 2000US-0235834P.	2000US-0234274P. 2000US-0234997P. 2000US-0234997P.	2000US-0233064P. 2000US-0233065P.	2000US-0232400P. 2000US-0232401P. 2000US-023262P	2000US-0232398P. 2000US-0232399P.	2000US-0231968P. 2000US-0232397P.	2000US-0232080P. 2000US-0232081P.	2000US-0231413P. 2000US-0231414P.	2000US-0231244P.	2000US-0231242P.	2000US-0230437P. 2000US-0230438P.	2000US-0229513P.	2000US-0229345P.	2000US-0229343F.	2000US-0229287P.	2000US-0227009P. 2000US-0228924P.	2000US-0227182P.	2000US-0226681P. 2000US-0226868P.	2000US-0225759P. 2000US-0226279P.	2000US-0225758P.	2000US-0225447P.	2000US-0225268P. 2000US-0225270P.	2000US-0225267P.	2000US-0225214P.	2000US-0224519P. 2000US-0225213P.	2000US-0224518P.	2000US-0220963P.	2000US-0217496P.	2000US-0217487P.	2000US-0216647P.	2000US-0215135P.	2000US-0209467P.	2000US-0198123P. 2000US-0205515P.	2000US-0190076P.	2000US-0186350P. 2000US-0189874P.	2000US-0184664P.	2000US-0179065P.	2001WO-US001354.	
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WPI; 2001-48	UMA-) HUN	-2000	08-DEC-2000;	2000	05-DEC-2000;	01-DEC-2000; 05-DEC-2000;	17-NOV-2000; 01-DEC-2000;	17-NOV-2000;	17-NOV-2000;	17-NOV-2000;	17-NOV-2000;	2000	-2000	-2000	2000	-2000	2000	-2000 -2000	-2000	2000	-2000	-2000	2000	-2000	-2000	2000	2000	2000	2000	2000	2000	-2000	-2000 -2000	-2000	-2000 -2000	2000		2000	SEP
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RESULT 47

AAK80185/c

ID

AAK80185;

XX

AC

AAK80185;

XX

AC

AAK80185;

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AC

AAK80185;

XX

AC

AAK80185;

XX

DT

O7-NOV-2001 (first entry)

XX

DE

Human immune/haematopoietic antigen

XX

Cytostatic; gene therapy; vaccine; immu

XX

Cytostatic; gene therapy; vaccine; immu

XX

OS

Homo sapiens.

XX

PM

W0200157182-A2.

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PM

W0200157182-A2.

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PM

APEB-2000; 2000US-0180628P.

PR

APAR-2000; 2000US-01806350P.

PR

APAR-2000; 2000US-020447P.

PR

APUL-2000; 2000US-021643P.

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APUL-2000; 2000US-021748P.

APUL-2000; 200
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Query Match
Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Matches 50;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Human immune/haematopoietic antigen genomic sequence SEQ ID NO:34997
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Sequence 13026 BP; 4098 A; 2489 C; 2384 G; 4055 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Disclosure; SEQ ID NO 34996; 3071pp + Sequence Listing; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  1.6%;
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Pred. No.
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o. 1.4e-12;
0;
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         14-SEP-2000

14-SEP-2000

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21-SEP-2000

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25-SEP-2000

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20-OCT-2000

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01-NOV-2000

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01-NOV-2000

01-NOV-2000
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22-AUG-2000;
22-AUG-2000;
23-AUG-2000;
01-SEP-2000;
01-SEP-2000;
01-SEP-2000;
01-SEP-2000;
05-SEP-2000;
05-SEP-2000;
06-SEP-2000;
06-SEP-2000;
08-SEP-2000;
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14-AUG-2000;
14-AUG-2000;
18-AUG-2000;
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14-AUG-2000;
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      2000US-022954P

2000US-0224519P

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2000US-022556PP

2000US-022526PP

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2000US-0225275PP

2000US-022575PP

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2000US-022668PP

2000US-022668PP

2000US-0229343PP

2000US-0229343PP

2000US-0229343PP

2000US-0239343PP

2000US-0231243PP

2000US-0231239PP

2000US-023123PP

2000U
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ARK54951 to ARK64702 encode the human immune/haematopoietic antigen (I) camino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic cc activity, and can be used in gene therapy and vaccine production. (I) cc proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For cc example, they may be used to treat disorders associated with decreased cc expression by rectifying mutations or deletions in a patient's genome cc that affect the activity of (I) by expressing inactive proteins or to cc supplement the patients own production of (I). Additionally, (I) cc polynucleotides may be used to produce the secreted (I), by inserting the cc mucleic acids into a host cell and culturing the cell to express the cc protein. (I) proteins and polynucleotides may be used to prevent, cc diagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metastases of haematopoietic active cells. AAK64703 cc AAK87694 represent human immune/haematopoietic antigen genomic
        08-NOV-2000
17-NOV-2000
17-NOV
                                                                                                                                                                                                                                                                                                                                                                                                                       Nucleic
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2000US-0249207P.
2000US-0249208P.
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English.

and metastasis.

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16-MAR-2000
18-ARR-2000
18-ARR-2000
19-MAY-2000
07-JUN-2000
28-JUN-2000
07-JUL-2000
07-JUL-2000
11-JUL-2000
11-JUL-2000
14-JUL-2000
14-AUG-2000
15-AUG-2000
16-AUG-2000
17-AUG-2000
18-AUG-2000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               21-NOV-2001 (first
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              AAL05461;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            sequences from the present invention. AAK54942 to AAK54950 and AAM82169 represent sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       17-JAN-2001;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCCAGAGACTCTGTCTC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 reproductive system related antigen; reproductive system disorder;
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2000US-0186628P

2000US-0186350P

2000US-0186350P

2000US-0198123P

2000US-0214867P

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2000US-022928P

2000US-022928P
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  1.6%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Score 50; DB; Pred. No. 1.4
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0;
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1.4e-12;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  G; 4055 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Length 13026;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 8149
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1-SEP-2000; 1-SEP-2000; 5-SEP-2000; 6-SEP-2000; 6-SEP-2000; 6-SEP-2000; 8-SEP-2000; 8-SEP-2000;

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RESULT 49
ABL98314/c
ID ABL983
XX ABL983
XX ABL983
XX ABL983
XX ABL983
XX Equation 21-JUN
XX Human;
XW reprod
XW reprod
XW cardio
XW gastro
XX Homo s
XX PN WO2001
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Best Local S
Matches 50
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01-DEC-2000
01-DEC-2000
05-DEC-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Isolated used in p
                                                          Human; testicular antigen; testes; cancer; metastasis; immune disorder; reproductive system disorder; urinary system disorder; gene therapy; cardiovascular disorder; respiratory disorder; neurological disorder; gastrointestinal disease; infection; cytostatic; gene; ds.
                                                                                                                                                                                                                                                                                                                                                                                      The present invention provides the protein and coding sequences of number of human reproductive system related antigens. These can be in the prevention and treatment of reproductive system disorders, including cancer. The present sequence is a genomic sequence encoding protein of the invention
                                                                                                                                                                                                ABL98314 standard; DNA; 31474
                                                                                                                                                                                                                                                                                                                                                                 Sequence
              WO200155317-A2
                                     Homo sapiens.
                                                                                                                                                21-JUN-2002
                                                                                                                                                                          ABL98314;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Disclosure;
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                                                                                                                                                                                                                                                                                                                Local Similarity
les 50; Conserv
                                                                                                                                                                                                                                                                                     3073
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                                                                                                                       testicular antigen
                                                                                                                                                                                                                                                               72
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        d nucleic acid molecule encoding a reproductive preventing, treating or ameliorating a medical
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                HUMAN
                                                                                                                                                                                                                                                                AGATTGTGCCACTGCAGCCTGGGCAACAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                31474 BP; 9245
                                                                                                                                                                                                                                                                                         AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Barash SC,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  SEQ ID NO 8149; 1297pp + Sequence Listing; English.
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2000US-0249215P.
2000US-0249214P.
2000US-0249214P.
2000US-0249245P.
2000US-0249245P.
2000US-0249264P.
2000US-0249264P.
2000US-0249269P.
2000US-0249299P.
2000US-0249299P.
2000US-0250160P.
2000US-0250160P.
2000US-02511030P.
2000US-02511030P.
2000US-02511868P.
2000US-02511868P.
2000US-02511869P.
2000US-02511869P.
2000US-0251989P.
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2000US-025199P.
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100.0%; Pr
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                                                                                                                                                                                                                                                                                                                                                                A;
                                                                                                                        encoding
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Pred. No.
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                                                                                                                                                                                                                                                                                                                 Mismatches
                                                                                                                        DNA
                                                                                                                                                                                                                                                                                                                                                                   6292
                                                                                                                        fragment
                                                                                                                                                                                                                                                                                                                            DB 4; Le
1.3e-12;
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01-SEP-2000
01-SEP-2000
01-SEP-2000
06-SEP-2000
06-SEP-2000
08-SEP-2000
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14-SEP-2000
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21-SEP-2000
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25-SEP-2000
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27-SEP-2000
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29-SEP-2000
29-SEP-2000
20-CCT-2000
20-CCT-2000
20-CCT-2000
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20-CCT-2000
20-CCT-2000
20-CCT-2000
20-CCT-2000
21-SEP-2000
21-NOV-2000
08-NOV-2000

2000US-022934AP.
2000US-0229549P.
2000US-023943P.
2000US-023124AP.
2000US-023124AP.
2000US-0231413P.
2000US-0231413P.
2000US-0231414P.
2000US-0231414P.
2000US-0232081P.
2000US-0232081P.
2000US-0232081P.
2000US-0232399P.
2000US-0232399P.
2000US-0232399P.
2000US-0232399P.
2000US-023363P.
2000US-0234274P.
2000US-0235834P.
2000US-0235834P.
2000US-0235834P.
2000US-0235834P.
2000US-024678P.
2000US-024678P.
2000US-024677P.
2000US-024677P.
2000US-0246528P.
2000US-0246528P.
2000US-0246528P.
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2000US-0246528P.
2000US-0246529P.
2000US-02465

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17-JAN-2000; 17-JAN-2000; 24-FEB-2000; 24-FEB-2000; 21-MAR-2000; 11-MAR-2000; 11-MAR-2000; 11-MAR-2000; 11-JUL-2000; 11-AUG-2000; 11-SEP-2000; 11-SE	- ATTG-
2001WO-US001329. 2000US-018065P. 2000US-0180668P. 2000US-0184664P. 2000US-018139P. 2000US-019133P. 2000US-02194867P. 2000US-02194869. 2000US-02194869. 2000US-02194899. 2000US-02194899. 2000US-02194899. 2000US-02194899. 2000US-0225213P. 2000US-0225266P. 2000US-0225266P. 2000US-0225266P. 2000US-0225266P. 2000US-02252709P. 2000US-0225270P. 2000US-0225347P. 2000US-0225349P. 2000US-0225349P. 2000US-022514P. 2000US-022514P. 2000US-022514P. 2000US-022514P. 2000US-022514P. 2000US-022514P. 2000US-022514P. 2000US-023141P. 2000US-02344P. 2000US-023	
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29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 20-CCT-2000; 02-CCT-2000; 02-CCT-2000; 13-CCT-2000; 20-CCT-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 20-NOV-2000; 21-NOV-2000; 21-NO	Ĺ
2000US-0233369 2000US-023369 2000US-023369 2000US-023369 2000US-0237039 2000US-0237039 2000US-0237039 2000US-024921 2000US-024921 2000US-0246477 2000US-0246478 2000US-0246528 2000US-0249218 2000US-025186 2000US-025186 2000US-025186 2000US-025186 2000US-0259678 CENOME SCI IN)US-0236368
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XFFX8X000000XX

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RESULT 50
AAS30115/c
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Best Local S
Matches 50
31-JAN-2000

04-FEB-2000

24-FEB-2000

02-MAR-2000

16-MAR-2000

11-MAR-2000

11-MAR-2000

11-MAY-2000

07-JUN-2000

07-JUN-2000

07-JUL-2000

07-JUL-2000

11-JUL-2000

11-JUL-2000
                                                                                                                                                                                                                                                                                                                                                                                                                               Lung antigen protein; human; mouse; rabbit; goat; horse; cat; dog; chicken; sheep; immunosuppressive; antiarthritic; vasotropic; antirheumatic; antiproliferative; cytostatic; cardiant; neuroprotective; cerebroprotective; nootropic; antibacterial; virucide; fungicide; cancer; ophthalmological; vulnerary; gene therapy; autoimmune disease; neoplasm; hyperproliferative disorder; breast; liver; cardiovascular disorder; ds; cerebrovascular disorder; nervous system disorder; bacterial infection; fungal infection; viral infection; ocular disorder; endocrine disorder; gastrointestinal disorder; renal disorder; respiratory disorder; wound healing; skin aging; organ transplantation; food preservative; tissue regeneration; anti-infertility; food additive.
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AAS30115
                                                                                                                                                                                                                                                                                                                                                                                                      Homo
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    The present invention provides the protein and coding sequences of 973 human testicular antigens, and fragments of their genomic sequences. The sequences can be used in the treatment of cardiovascular, urinary system, reproductive system, immune, respiratory, neurological and gastrointestinal disorders, infections, and particularly cancer, especially testicular cancers. The present sequence is a DNA encoding a
                                                                                                                                                                                                                                                                                                                                        02-AUG-2001
                                                                                                                                                                                                                                                                                                                                                                      WO200155303-A2
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Human
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                21-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               AAS30115;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 31474 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Disclosure; SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Nucleic acids encoding 973 human testicular antigen polypeptides, useful for preventing, diagnosing and/or treating testicular cancer.
                                                                                                                                                                                                                                                                                                       17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           3073
                                                                                                                                                                                                                                                                                                                                                                                                    sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              lung
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             72
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             50;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        fragment of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 23
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              antigen genomic DNA #185.
2000US-0179065P.
2000US-0184664P.
2000US-0186350P.
2000US-0186350P.
2000US-0198074P.
2000US-0198123P.
2000US-0205515P.
2000US-0214866P.
2000US-0215135P.
2000US-0215135P.
2000US-0216647P.
2000US-021684P.
2000US-0217487P.
2000US-0217487P.
2000US-0217486P.
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2000US-0217487P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DNA;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         1.6%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               32189
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Score 50; DB
Pred. No. 1.3
0; Mismatches
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J. 1.3e-12;
O;
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   14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
11-AUG-2000
12-AUG-2000
22-AUG-2000
22-AUG-2000
23-AUG-2000
01-SEP-2000
02-CCT-2000
03-CCT-2000
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14-AUG-2000;
2000US-0224518P.
2000US-0225266P.
2000US-0225266P.
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2000US-0241284P.
2000US-0241284P.
2000US-0241284P.
2000US-0241849P.
2000US-0241849P.
2000US-0244617P.
2000US-0246479P.
2000US-0246479P.
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Sequences AAS2991-AAS30164 represent genomic DNA molecules, which encode the lung antigen polypeptides of the invention. Lung antigen polypeptides and their associated polymucleotides are useful in the diagnosis, treatment and prevention of various types of disorders in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. A pathological condition can be determined by detecting the presence or absence of a mutation in a lung antigen polynucleotide. The treatable disorders include autoimmune diseases such as rheumatoid arthritis, hyperproliferative disorders such as neoplasms of the breast or liver, cardiovascular disorders such as cardiac arrest, cerebrovascular disorders such as cardiac arrest, cerebrovascular disorders such as cardiac arrest, cerebrovascular of liver, cardiovascular disorders such as cardiac arrest, viruses and fungi, ocular disorders such as corneal infection, endocrine disorders such as premature labour and infertility, gastrointestinal disorders such as corneal alignyders such as corneative and such as corneative and an endocrine such as premature labour and infertility, gastrointestinal disorders such as
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08-NOV-2000
08-NOV-2000
17-NOV-2000
17-NOV
premature
Crohn's di
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                                                                                                                                                                                                                                                                                                                                                             Claim 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                      Isolated
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (HUMA-) HUMAN GENOME SCI INC
                                                                                                                                                                                                                                                                                                                                                                                                   testing and detection e.g. diagnosis.
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                                                                                                                                                                                                                                                                                                                                                                                                                            polypeptide for treating, preventing and/ or prognosing ory disorders related to the lung including lung cancers
                                                                                                                                                                                                                                                                                                                                                             SEQ ID NO 379; 507pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Barash
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2000US-0249214P.
2000US-0249216P.
2000US-0249214P.
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2000US-0246524P.
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2000US-0246611P.
2000US-0246611P.
2000US-0246611P.
2000US-0246611P.
2000US-024611P.
2000US-024921P.
2000US-024921P.
2000US-024921P.
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such as glomerulonephritis
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RESULT 51
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24-FBB-2000
02-MAR-2000
16-MAR-2000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        gene therapy; lung antigen; neoplasia; acute myelogenous leukaemia; adenocarcinoma; respiratory disorder; chronic rhinitis; sinusitis; immunodeficiency; X-linked agammaglobulinaemia; X-linked infantile agammaglobulinaemia; inflammatory disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human novel lung related polypeptide DNA SEQ
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llarity 100.0%; P
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The invention relates to an isolated lung antigen polypeptide sequence or composed sequence in a cDNA clone. The polypeptide and its polynucleotide are useful for treating, preventing, diagnosing and/or prognosing concepts as a cutte myelogenous leukaemias, adenocarcinoma; respiratory consoplasias e.g. acute myelogenous leukaemias, adenocarcinoma; respiratory consoplasias e.g. acute myelogenous leukaemias, adenocarcinoma; respiratory consoplasias e.g. acute myelogenous leukaemias; immunodeficiencies such as chronic rhinitis, sinusitis; immunodeficiencies such as chronic rhinitis, alvoslitis; immunodeficiencies such as chromatory disorders such as adrenalitis, alvoslitis; immune complex configuratory disorders such as serum sickness, polyarteritis nodosa; bleeding disorders concertion such as kidney failure, multiple myeloma; disorders consociated with macrophage numbers and/or macrophage function such as colon cancer; caucher's disease, Neimann-Pick disease; tumours such as colon cancer; consociated with as Albers-Schonberg disease, bowlegs; muscle consociated such as as a schone is schone in traumatic lesions.
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standard;
(first entry)
                           DNA;
                            32193
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musculoskeletal system related polynucleotide SEQ Ħ ð

Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral; antiallergic; hepatotropic; antidiabetic; antiinflammatory; antiulcer; vulnerary; anticonvulsant; antibacterial; antifungal; antiparasitic; cardiant; gene therapy; cancer; immune disorder; cardiovascular disorder; neurological disease; infection; human; secreted protein; musculoskeletal system; ds.

2000US-0179065P.
2000US-018664P.
2000US-0184664P.
2000US-0184664P.
2000US-0199076P.
2000US-0199076P.
2000US-0199076P.
2000US-0214886P.
2000US-0216479P.
2000US-0216479P.
2000US-0217496P.
2000US-0217496P.
2000US-0217496P.
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2000US-0225214P.
2000US-0225214P.
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2000US-0229343P.
2000US-023943P.
2000US-023943P. 2001WO-US001338

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2000US-024677P
2000US-0241785P
2000US-024677P
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CC preventing, treating or ameliorating medical conditions e.g. by protein cc or gene therapy. The genes are isolated for a ready and protein cc and (ant) agonists are useful in the diagnosis, treatment and prevention cf (a) cancer, e.g. breast and ovarian cancer and other cancers of the cd llurg, or urogenital; (b) immune disorders e.g. Addison's disease, cc allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis, cc diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid cc arthritis and ulcerative colitis; (c) cardiovascular disorders such as cc myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and epilepsy; and (f) infectious diseases such as vital, fungal and parasitic infections. Note: The sequence data for obtained in electronic format directly from WIPO at (c) cardiovascular disorders diseases obtained in electronic format directly from WIPO at (c) cardiovascular disorders such as concernat did not form part of the printed specification, but was obtained in electronic format directly from WIPO at
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
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17-NOV-2000;
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08-DEC-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Isolated polypeptide disorders related to cancers and also for
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          for treating, preventing and/or prognosing the musculoskeletal system including muscultesting and detection e.g. diagnosis.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Ruben
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musculoskeletal

RESULT 54
ABX59246/c
ID ABX592
XX
AC ABX592
XY
AC ABX592
XY
DT 26-FEB
XX
DE CDNA e
XX
Gene; 밁 გ Matches Query Match Best Local (Gene; ss; musculoskeletal system antigen; cancer; metastasis; cDNA encoding novel human musculoskeletal system antigen #1590. ABX59246 standard; Sequence 32193 BP; 26-FEB-2003 28855 1.6%; { 1 Similarity 100.0%; 50; Conservative 0; (first entry) CDNA; 32193 10182 A; Score 50; DB; Pred. No. 1.3 6701 C; ВP 6066 1.3e-G; 9244 T; 0 U; 0 Other; 4 0 Length 32193; Indels <u>,</u> Gaps 0

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     post-operative tissue repair; limb regeneration; neuronal growth; neurodegenerative disorder; Alzheimer's disease; Parkinson's disease; AIDS-related complex; chondrocyte growth; bone regeneration;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           cardiovascular condition; wound; post-operative tissue repair; li
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           10-OCT-2002
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2000US-0220964P.
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RESULT 55
ADG62943/c
ID ADG629
XX
AC ADG629
XX
DT 11-MAR
XX
DE Genomi
XX
KW neurop
KW gastro

(first entry)

ADG62943 standard; DNA; 32193 BP

Genomic DNA 11-MAR-2004 ADG62943

encoding human NOVX protein seg id

37

neuroprotective; nootropic; respiratory; cardiovascular;
gastrointestinal; antiparkinsonian; immunosuppressive; dermatological;

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AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 28806 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122 Matches

Local

Similarity

Conservative

<u>,</u>

Pred. No. Mismatches

1.3e-12 0

Indels

0;

Gaps

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CC and limb regeneration; stimulates neuronal growth; can treat and prevent CC neuronal damage occurring in certain disease, Parkinson's disease, and AIDS-CC conditions, such as, Alzheimer's disease, Parkinson's disease, and AIDS-CC related complex; stimulates chondrocyte growth, thus they can be used to cenhance bone and periodontal regeneration and aid in tissue transports or CC bone grafts; prevents skin aging due to sumburn by stimulating cC enthance bone and periodontal regeneration and aid in tissue transports or CC bone grafts; prevents skin aging due to sumburn by stimulating cC enthance bone and periodotes melanocyte growth; stimulates or comports or comports or comports of the sumburn by stimulating cC growth and differentiation of hematopoietic cells and bone marrow cells when used in combination with other cytokines; maintains organs before comports or decreases the differentiation of primary tissues; cincreases or decreases the differentiation or proliferation of embryos; cc increases or decreases the differentiation or proliferation of embryos; cc increases or decreases bedy height, weight, hair colour, eye colour, cased in percentage of adipose tissue, pigmentation, size, and shape (e.g., cc skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cc pabilities, hormonal or endocrine levels, appetite, libido, memory, or creases; increases or decreases storage capabilities, fat content, lipid, components. This sequence encodes a novel human musculoskeletal system cc printed specification, but was obtained in electronic format directly
                   Query Match
                                                                                                       printed specification, but was obtained in electronic from the US patent office at ftp.seqdata.uspto.gov/sequence.html?DocID=20020147140
                                                                 Sequence 32193 BP; 10182 A; 6701 C; 6066 G; 9244 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   tissues associated with conditions such as thrombosis, arteriosclerosis, and other cardiovascular conditions; treats wounds due to injuries,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   sequence encoding musculoskeletal system associated polypeptides useful for detecting disorders, e.g., cancer or cancer metastases, in animals humans. The nucleic acid: stimulates re-vascularisation of ischaemic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Disclosure; SEQ ID NO 2623; 321pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Rosen CA, Ruben SM,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     (ROSE/)
(RUBE/)
(BARA/)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The invention describes an isolated nucleic acid
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          associated
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Isolated nucleic acid
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              08-DEC-2000; 2000US-0251869P.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  post-operative tissue repair, and ulcers; stimulates angiogenesis
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        RUBEN S
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     BARASH S C
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ucleic acid molecules encoding musculoskeletal polypeptides, useful for detecting disorders,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          3 >
1.6%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Barash
                                                                                                                                                             but was obtained in electronic format directly
                   Score 50;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         ၁၁
                     DB 8;
                   Length 32193;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            molecule comprising
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   system
e.g. cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           in animals or
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artialrimamatory, attirhoumatic, antiatrico, antiatraemic) artialrimamatory, attirhoumatic, antiatrico, antiatraemic) artialrication colored c
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2000US-0232397P. 2000US-0232398P. 2000US-0232399P. 2000US-0232401P. 2000US-0233063P. 2000US-0233063P. 2000US-0233063P. 2000US-0233063P. 2000US-023423P. 2000US-023427P. 2000US-023427P. 2000US-023637P. 2000US-02496602P. 2000US-0249669P. 2000US-0249667P. 2000US-0246611P. 2000US-024661P. 2000US-024661P. 2000US-024661P. 2000US-024661P. 2000US-024921P. 2000US-024920P. 2000US-025039P. 2000US-025039P.

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ADJ29996/c
ID ADJ299
XX
AC ADJ299
AC ADJ299
XX
DE Human
XX
KW muscul
KW gene t
XX
OS Homo s
XX
PN US2004
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PN US2004
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XX
NS 15-JAN
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              CC The invention describes an isolated nucleic acid molecule (I) encoding a CC protein comprising a sequence that is at least 95% identical to: a CC polynucleotide fragment of any of the nucleotide sequences listed in the CC specification, or of the cDNA sequences listed in the specification, combined in the specification of the cDNA sequences listed in the specification, combined and sequences listed in the specification, or a ccoding a polypeptide or a polypeptide fragment, domain or epitope of CC any of the amino acid sequences listed in the specification, or a ccode above, a polynucleotide which is an (allelic) combined above, a polynucleotide which is an (allelic) combined above, a polynucleotide which is an (allelic) combined above, a polynucleotide which is an extringent conditions to any of the above polynucleotides, where the polynucleotide does not hybridise under stringent conditions to a nucleic acid molecule and polypeptide are useful in diagnosing, preventing, prognosing cor treating diseases or disorders associated with aberrant expression candor activity of the above polypeptide, such as neural disorders, immune system disorders, muscular disorders, reproductive disorders.
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Best Local Similarity
Matches 50; Conserv
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                          15-JAN-2004
                                                                 US2004009488-A1
                                                                                                                                             musculoskeletal system; cytostatic; osteopathic; cancer; osteoporosis; gene therapy; vaccine; human; ds.
                                                                                                                                                                                                          Human musculoskeletal system-associated genomic DNA - SEQ ID 2623
                                                                                                                                                                                                                                                        20-MAY-2004
                                                                                                                                                                                                                                                                                                 ADJ29996;
                                                                                                                                                                                                                                                                                                                                      ADJ29996 standard; DNA; 32193 BP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             treating dispolypeptide,
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                                                                                                                                                                                                                                                                                                                                                                                                                                              28855
                                                                                                                                                                                                                                                                                                                                                                                                                                        polypeptides and nucleic acid molecules for diagnosing, preventing on atting diseases associated with aberrant expression or activity of the ypeptide, e.g. cancer, asthma, AIDS, Parkinson's disease or diabetes.
                                                                                                        sapiens
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llarity 100.0%;
Conservative (
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2000US-0256719P.

2000US-0251479P.

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2000US-0251868P.

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2000US-0251990P.

2000US-0254097P.

2001US-0259678P.

2001US-0259678P.

2001US-00764883.
                                                                                                                                                                                                                                                   (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Barash
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               0
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Score 50;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           SC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      DB 10;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1.3e-12;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               0
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Length 32193;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Gaps
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             0
    $\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac{1}{2}$\frac
  14-AUG-2000
118-AUG-2000
22-AUG-2000
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23-AUG-2000
30-AUG-2000
01-SEP-2000
01-SEP-2000
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25-SEP-2000
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25-SEP-2000
25-SEP-2000
29-SEP-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         11-JUL-2000;
11-JUL-2000;
14-JUL-2000;
26-JUL-2000;
26-JUL-2000;
14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         31-JAN-2000;
04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAY-2000;
07-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         30-JUN-2000;
07-JUL-2000;
07-JUL-2000;
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14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 14-AUG-2000
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2000US-018464P.
2000US-0189374P.
2000US-0189374P.
2000US-0199173P.
2000US-020515P.
2000US-0215135P.
2000US-0215135P.
2000US-0215135P.
2000US-0215135P.
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2000US-0217487P.
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2000US-0225214P.
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2000US-0225214P.
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2000US-022575P.
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2000US-022575P.
2000US-0231414P.
2000US-0231444P.
2000US-0231442P.
2000US-0231444P.
2000US-02314
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             2000US-0180628P.
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29-SBP-2000
02-OCT-2000
02-OCT-2000
02-OCT-2000
02-OCT-2000
03-OCT-2000
03-OCT-2000
03-OCT-2000
03-OCT-2000
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03-NOV-2000
08-NOV-2000
09-DC-2000
01-NOV-2000
01-NOV-2000
01-DC-2000
01-NOV-2000
01-DC-2000
                              Rosen CA,
2004-090458/09
                                                              HUMAN
                              Ruben SM,
                                                                                                                                                                                                                          2000US-0240960P
2000US-0241785P
2000US-0241786P
2000US-0241808P
2000US-0241809P
2000US-02446178P
2000US-0246475P
2000US-02464778P
2000US-0246477P
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2000US-0246525P
2000US-0246525P
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2000US-0246510P
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2000US-0259390P
2000US-0259391P
2000US-02567198P
                                                                                             2000US-0251990P.
2000US-0254097P.
2001US-0259678P.
2001US-00764877.
                                                                                                                                                           2000US-0251856P.
2000US-0251868P.
2000US-0251869P.
2000US-0251989P.
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                                                              GENOME
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ק הקרק הרואה ה הרואה הר

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FFFXSXSSSSSSSSSXX
                                                                                                                           Disclosure; SEQ ID NO 2623; 289pp; English.
                                                                                                                                               preventing, treamusculoskeletal
                                                                                                                                                         nucleic acid molecule, useful for preparing a medicament for venting, treating or ameliorating a medical condition e.g., cancer
                                                                                                                                                tissues or osteoporosis.
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Fi
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The invention relates to a novel isolated musculoskeletal system-associated nucleic acid molecule. The nucleic acid of the invention demonstrates cytostatic and osteopathic activities and may be useful for preparing a medicament for preventing, treating or ameliorating a medical condition such as cancer of the musculoskeletal tissues or osteoporosis, possibly via gene therapy or vaccine production. The current sequence is that of the human musculoskeletal system-associated genomic DNA of the invention. The current sequence is not shown within the specification per se but is available on the USPTO web-site http:seqdata.uspto.gov/sequence.html?DocID=20040009488.

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Query Match
Best Local S
Matches 50
                                   Sequence
50;
         Similarity
                                   32193 BP; 10182 A; 6701 C; 6066 G; 9244 T; 0 U;
Conservative
        1.6%;
0;
        Score 50;
Pred. No.
 Mismatches
           DB 12;
1.3e-12
         .3e-12;
                  Length
 Indels
                                     0 Other;
0
Gaps
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3122

0

3073

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RESULT 57
AAS30113/c
吊
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                                                                                                                                                                                                                                                                                                                                                             Lung antigen protein; human; mouse; rabbit; goat; horse; cat; dog; chicken; sheep; immunosuppressive; antiarthritic; vasotropic; antirheumatic; antiproliferative; cytostatic; cardiant; neuroprotective; cerebroprotective; nootropic; antibacterial; virucide; fungicide; cancer; ophthalmological; vulnerary; gene therapy; autoimmune disease; neoplasm; hyperproliferative disorder; brasst; liver; cardiovascular disorder; derevous system disorder; bacterial infection; cerebrovascular disorder; responsation; autoimmune disease; liver; cardiovascular disorder; despression; system disorder; bacterial infection; fungal infection; ocular disorder; endocrine disorder; gastrointestinal disorder; renal disorder; respiratory disorder; wound healing; skin aging; organ transplantation; food preservative; tissue regeneration; anti-infertility; food additive.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             AAS30113
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 21-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               AAS30113;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human lung antigen genomic DNA #183.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       28855
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               standard,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             32221
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31-JAN-2000 04-FEB-2000 24-FEB-2000 02-MAR-2000 11-MAR-2000 11-MAR-2000 18-APR-2000 07-JUN-2000 28-JUN-2000 28-JUN-2000 30-JUN-2000 07-JUL-2000 07-JUL-2000 11-JUL-2000 11-JUL-2000 11-JUL-2000 11-JUL-2000

2000US-0186350P. 2000US-0189874P.

2000US-0184664P

2000US-0190076P.
2000US-0205515P.
2000US-0205515P.
2000US-0209467P.
2000US-0215135P.
2000US-0216647P.
2000US-0216647P.
2000US-0217488PP.
2000US-0217486P.

Homo sapiens.

17-JAN-2001; 02-AUG-2001. WO200155303-A2

2001WO-US001301.

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08-NOV-2000
17-NOV-2000
17-NOV
Sequences AAS29931-AAS30164 represent genomic DNA molecules, which ence the lung antigen polypeptides of the invention. Lung antigen polypeptil and their associated polynucleotides are useful in the diagnosis, treatment and prevention of various types of disorders in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. A pathological condition can be determined by detecting the presence or absence of a mutation in a lung antigen polynucleotide. The treatable disorders include autoimmune diseases such as rheumatoid arthritis, hyperproliferative disorders such as neoplasms of the breast or liver, cardiovascular disorders such as cardiac arrest, cerebrovascular disorders such as cardiac arrest, cerebrovascular disorders such as cerebral ischaemia, nervous system disorders such as Alzheimer's disease, infections caused by bacteria, viruses and fungi,
                                                                                                                                                                                                                                                                                            Isolated polypeptide for treating, preventing and/or prognosing respiratory disorders related to the lung including lung cancers for testing and detection e.g. diagnosis.
                                                                                                                                                                                                                                                           Claim
                                                                                                                                                                                                                                                                                                                                                                                                                        Rogen
                                                                                                                                                                                                                                                                                                                                                                                                                                                             (HUMA-)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                               HUMAN
                                                                                                                                                                                                                                                           SEQ
                                                                                                                                                                                                                                                                                                                                                                                                                      Barash
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  2000US-0246474P.
2000US-0246475P.
2000US-0246475P.
2000US-0246524P.
2000US-0246525P.
2000US-0246528P.
2000US-0246610P.
2000US-0246610P.
2000US-0246611P.
2000US-0246611P.
2000US-0249210P.
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2000US-0249211P.
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2000US-0249219P.
2000US-0249219P.
2000US-0251988P.
2000US-0251868P.
2000US-0251868P.
2000US-0251868P.
2000US-0251869P.
2000US-0251989P.
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                                                                                                                                                                                                                                                                                                                                                                                                                      Ruben
                                                                                                                                                                                                                                                         507pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                        Z.
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which encode polypeptides

and also

14-JUL-2000
26-JUL-2000
14-AUG-2000
11-AUG-2000
11-AUG

2000US-0218290P.
2000US-022954P.
2000US-0225266P.
2000US-0225266P.
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2000US-0225279P.
2000US-0225758P.
2000US-0235897P.
2000US-0231413P.
2000US-

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RESULT 58
ADB33450 c
ID
ADB33450 standard; DNA; 32221 BP
XX
AC
ADB33450;
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C
ADB33450;
XX
C
ADB33450;
XX
DE
O4-DEC-2003 (first entry)
XX
DE
Human novel lung related polypep
XX
DE
Human novel lung antigen; neop
XX
DE
Human novel lung antigen; neop
XX
DE
XX
DE
XX
DE
XX
Von Willebrand's disease; acquir
XX
DECHANA-1003
XX
DECHANA-2003
XX
DECHANA-2003
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D2-MAR-2000; 2000US-0189133P
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D3-JUN-2000; 2000US-0189133P
PR
D3-JUN-2000; 2000US-029467P
PR
D3-JUN-2000; 2000US-021686P
PR
D3-JUN-2000; 200US-02168F
PR

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Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    ocular disorders such as corneal infection, endocrine disorders such as premature labour and infertility, gastrointestinal disorders such as Crohn's disease, renal disorders such as glomerulonephritis and respiratory disorders such as asthma and pleurisy. The polypeptides can also be used to aid wound healing, to prevent skin aging due to sunburn, to maintain organs before transplantation, to regenerate tissues and in chemotaxis. The polypeptides can also be used as a food additive or preservative to increase or decrease storage capabilities. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published pct sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             immunodeficiency; X linked agammaglobulinaemia;
X-linked infantile agammaglobulinaemia; inflammatory disorder;
Actinked infantile agammaglobulinaemia; inflammatory disorder;
adrenalitis; alveolitis; immune complex disease; serum sickness;
polyarteritis nodosa; bleeding disorder; thrombocytopenia;
Von Willebrand's disease; acquired platelet dysfunction; kidney fail
multiple mysloma, macrophage related disorder; daucher's disease;
Neimann-Pick disease; tumour; colon cancer; pancreatic cancer;
renal disorder; nephritis; bone disorder; Albers-Schonberg disease;
bowleg; muscle disorder; Becker's muscular dystrophy;
buchenne's muscular dystrophy; nervous disorder; ischaemic lesion;
traumatic lesion; endocrine disorder; Cushing's syndrome;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                gene therapy; lung antigen; neoplasia; acute myelogenous leukaemia; adenocarrinoma; respiratory disorder; chronic rhinitis; sinusitis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Human novel lung related polypeptide DNA SEQ ID NO 377.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               1.6%;
ilarity 100.0%;
Conservative (
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             deficiency; gastrointestinal
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ; Score 50; DB
i; Pred. No. 1.3
0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      DB 5; 4...
3. 1.3e-12;
0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             disorder; dysphagia;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Length 32221;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       kidney failure;
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11-SEP-2000
11-SEP
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17-NOV-2000

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17-NOV-2000
The invention relates to an isolated lung antigen polypeptide sequence or encoded sequence in a cDNA clone. The polypeptide and its polynucleotide are useful for treating, preventing, diagnosing and/or prognosing diseases and/or disorders such as pathological cell proliferative neoplasias e.g. acute myelogenous leukaemias, adenocarcinoma; respiratory disorders such as chronic rhinitis, sinusitis; immunodeficiencies such as x-linked agammaglobulinaemia; inflammatory disorders such as drenalitis, alveolitis; immune complex diseases such as serum sickness, polyarteritis nodosa; bleeding disorders such as thrombocytopenia, von Willebrand's disease; acquired platelet dysfunction such as kidney failure, multiple myeloma; disorders associated with macrophage numbers and/or macrophage function such as Gaucher's disease, Neimann-Pick disease; tumours such as colon cancer, pancreatic cancer; renal disorders such as kidney failure, nephritis;
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17-NOV-2000;
17-NOV-2000;
                                                                                                                                                                                                                                diagnosing acute
Von Willebrand's
                                                                                                                                                                                                                                                                                         WPI;
                                                                                                                                                                                                       Disclosure; SEQ ID NO 377; 178pp; English.
                                                                                                                                                                                                                                Novel isolated lung antigen polypeptides useful for treating, preventing, diagnosing acute myelogenous leukemias, adenocarcinoma, thrombocytopenia, von Willebrand's disease.
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17-NOV-2000;
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                                                                                                                                                                                       The invention relates to a method for identifying a molecule involved in lipid regulation comprising identifying a molecule that binds to or inhibits binding of a molecule to high bone mass (HBM) or its wild type gene, Zmax1. Compounds identified by the method are useful for treating, diagnosing, preventing or screening for normal and abnormal lipid-associated conditions, including arterioscleris, cardiovascular disease, stroke, and osteoporosis. The compounds may also be used in the treatment or prevention of diabetic atherosclerosis, neurovascular conditions caused by plaque build-up, poor circulation due to plaque build-up and associated poor wound-healing. The methods may be used in gene therapy, pharmaceutical development, and diagnostic assays for bone development disorders. Molecules identified by comparison of Zmax1 and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Identifying molecules involved in lipid regulation, useful for diagnosing, treating or preventing e.g., arteriosclerosis, comgidentifying a molecule that binds to high bone mass gene or its corresponding wild type gene.
                                                                                        HBM systems can be used as surrogate markers in pharmaceutical development, in diagnosis of human or animal bone disease, and in treatment of bone diseases. Sequences ABK23776-ABK33411 represent
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   lipid-associated condition; arteriosclerosis; cardiovascular disease; ss; osteoporosis; atherosclerosis; diabetic atherosclerosis; plaque build-up; neurovascular condition; wound healing; gene therapy; PCR primer; probe;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  bone disorders such as Albers-Schonberg disease, bowlegs; muscle disorders such as Becker's muscular dystrophy, Duchenne's muscular dystrophy; nervous disorders such as ischaemic lesions, traumatic lesions; endocrine disorders such as Cushing's syndrome, corticosteroid
                                     molecules encoding human Zmax1 and HBM, and and adapters of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Carulli JP,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      osteopathic;
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Similarity 100.0%;
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              English
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                                                                PCR primers,
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                                                                                                                                                                                                                                                                         The present invention relates to novel DNA and protein sequences which care associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bioactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bioactive agent capable of modulating the activity of CAP; (iv) for carcinoma; (vi) for inhibiting the activity of CAP; (vi) for treating carcinoma; (vii) for neutralizing the effect of CAP; (vii) for treating carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip; (cx) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent
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ACN44786 standard; DNA; 156843 BP.
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                                                                                                                                                                                                                                                                         Sequence 66973
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          comprises a nucleotide sequence.
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                                                                                                                                               3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
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                                                                                                                                                 1.6%; suc. 100.0%; Pred. No. 1
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Pred. No. 1.3e-12;
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hes 0;
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                                                                                                                                                                                                                             Length 66973;
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RESULT 62
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  Homo sapiens.
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Query Match
Best Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   determining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent US2002182586A1, for which no sequence data was published
Human; Chromosome 5; ds; gene; ADAM19; Endophilin 1; Endophilin 2; NRG2; ADAMTS2; a disintegrin and metalloprotease; neuroregulin 2; SNP; single nucleotide polymorphism; a disintegrin and metalloprotease with thrombospondin type1 motif 2; a sthma; atopy; obesity; inflammatory bowel disease; respiratory disorder.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Recombinant nucleic acid useful for diagnosis and treatment comprises a nucleotide sequence.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Human genomic sequence hCG27192
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Sequence 156843 BP; 33001 A; 41006 C; 43823 G; 38715 T; 0 U; 298 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Claim 1; SEQ ID NO 1408; Opp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           WPI; 2003-328604/31.
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                                                                                                                                                                                                        Human ADAMTS2 gene.
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Pred. No. 1.2e-12;
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Single nucleotide polymorphism*	/*tag= av /standard_name= "Single		FT	97746,T)	variation
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ngle nucleotide polymorphism" Figuration Fi	replace (215294, A) /*tag= am	variation	1 P P P	ard name= "Single nucleotide e(170372,G)	variation
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ngle nucleotide polymorphism" FT variation FT ration FT ration FT variation	replace (213555, A) /*tag= aj /standard name= "Sing	<u>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</u>		"Single nucleotide A)	variation
ngle nucleotide polymorphism" FT variation FT ration FT ration FT ration FT ration FT ration FT ration FT ratiation	replace (213324, A) /*tag= ai /standard name= "Sing	variation		_name= "Single nucleotide 43748,T)	variation
ngle nucleotide polymorphism" FT variation FT ration FT variation	replace (213294, A) /*tag= ah /standard name= "Sing	variation	리 다 다 :	Single nucleotide	variation
ngle nucleotide polymorphism" FT variation FT FT variation	/standard_name= "Sing replace(213243,A) /*tag= ag /standard_name= "Sing	variation	ים ען די יים על על די	Single nucleotide	variation
ngle nucleotide polymorphism" FT variation FT PT PT variation FT PT	/standard_name= "Sing replace(211462,T) /*tag= af	variation	FT	Single nucleotide	variation
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ngle nucleotide polymorphism" refrection for the polymorphism for the p	<pre>/standard_name= "Sing replace(211213,A) /*tag= ad</pre>		r r r	e ard_name= "Single nucleotide a(143591,G)	variation
ngle nucleotide polymorphism" refr variation refr variation refrection	/standard_name= "Sing replace(199313,C) /*tag= ac	variation	FT FT	Single nucleotide	e t
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RESULT
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Matches 50
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17204/c
AAK77204 standard; DNA; 95 BP.
31-JAN-2000;
04-FEB-2000;
24-FEB-2000;
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Allen K;
                                                                                                                                                                                                                 Human; immune; haematopoietic; immune/haematopoietic cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               The invention relates
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Claim 2; SEQ ID NO 9; 338pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 New isolated nucleic acid or alternate splice variant, useful for diagnosing and treating a disintegrin and metalloprotease (ADAM) or interactor gene-associated disorder, e.g. asthma, atopy, obesity or inflammatory bowel disease.
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                                                                        17-JAN-2001; 2001WO-US001354.
                                                                                                           09-AUG-2001.
                                                                                                                                          WO200157182-A2
                                                                                                                                                                                 Homo
                                                                                                                                                                                                                                                                     Human immune/haematopoietic antigen genomic sequence SEQ ID NO:32016.
                                                                                                                                                                                                                                                                                                        07-NOV-2001
                                                                                                                                                                                                                                                                                                                                           AAK77204;
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2000US-0179065P.
2000US-0180628P.
2000US-0184664P.
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2000US-0199074P.
2000US-01990759.
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2000US-029467P.
2000US-0216487P.
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2000US-0225747P.
2000US-0225758P.
2000US-0225758P.
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08-NOV-2000
AAK54951 to AAK64702 encode the human immune/haematopoietic antigen
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                        Disclosure;
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17-NOV-2000;
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                                                for
                                               acids encoding human immune/hematopoietic antigen polypeptides, for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                             Barash SC,
                                                                                                                                                          2000US-0246524P.
2000US-0246525P.
2000US-0246528P.
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2000US-0246611P.
2000US-0249201P.
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2000US-0246478P
                        SEQ ID NO 32016; 3071pp + Sequence Listing; English.
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2000US-0239935P.
                                                                                                                                     GENOME SCI INC.
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RESULT 64
AAK77203/c
ID AAK77203/c
XX AAK77203;
XX AAK77203;
XX AAK77203;
XX AAK77203;
XX AAK77203;
XX Human immu
XX Human; imm
XW Cytostatic
XX Homo sapic
XX II-JAN-200
PF 17-JAN-200
PF 17-JAN-200
PR 16-MAR-200
PR 16-MAR-200
PR 16-MAR-200
PR 19-MAY-200
PR 19-MAY-200
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Matches 49; Conserv
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                      24-FBB-2000
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04-FEB-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   09-AUG-2001
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Pred. No.
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Rosen ξ Barash SC, XX.

2001-483426/52

Nucleic useful for for preventing, acids encoding human immune/hematopoietic antigen polypeptides, for preventing, diagnosing and/or treating cancers and metastasis.

Disclosure; SEQ ID ŏ 32015; 3071pp + Sequence Listing; English.

CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic contivity, and can be used in gene therapy and vaccine production. (I) crotesins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For ceample, they may be used to treat disporders associated with decreased cexpression by rectifying mutations or deletions in a patient's genome cc that affect the activity of (I) by expressing inactive proteins or to concert affect the patients own production of (I). Additionally, (I) colored colo

Sequence 95 B₽; 22 A; 14 C; 16 G; 4 H 0 U; 0 Other;

Query Match Best Local Similarity 1.6%; Score 49; DB 4; L Pred. No. 4.7e-12; Length

NAK86736 standard; DNA; 272 BP. AAK86736; 07-NOV-2001 (first entry) Ruman immune/haematopoietic antigen genomic sequence SEQ ID Human; immune/haematopoietic; immune/haematopoietic antige cytostatic; gene therapy; vaccine; metastasis; ds. Homo sapiens. W0200157182-A2. 09-AUG-2001. 17-JAN-2001; 2001WS-019062B, 4-EBB-2000; 2000US-018625B, 4-EBB-2000; 2000US-018635B, 4-EBB-2000; 2000US-01	Matches 49; Conservative 0; Mismatches 0; Indels 0; Gaps 0; Qy 3073 AGATTGTGCCACTGCAGCCTGGGCAACAGAGCAAGACTCTGTCT 3121

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02333 02334 02344 02444 02446 0246 0246 0246 0246 0246	000US-02312/ 000US-02312/ 000US-02314/ 000US-02314/ 000US-02320/

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ARSSULT 66
AAXS6737/c
ID AAXS677
XX AAXS677
XX AAXS677
XX O7-NOV
DT 07-NOV
XX Y
XX Human
XX Human;
XW Human;
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                                                                                                                                                                                                                                                                                                                                                                                                                                  CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic contivity, and can be used in gene therapy and vaccine production. (I) cc proteins and polynucleotides may be used in the prevention, diagnosis and ct treatment of diseases associated with inappropriate (I) expression. For cc example, they may be used to treat disorders associated with decreased cexpression by rectifying mutations or deletions in a patient's genome ct that affect the activity of (I) by expressing inactive proteins or to complement the patients own production of (I). Additionally, (I) cc polynucleotides may be used to produce the secreted (I), by inserting the cc protein. (I) proteins and polynucleotides may be used to prevent, cc diagnose and treat immune/haematopoietic-related diseases, especially cc cancers and cancer metastases of haematopoietic-derived cells. AAK64703 cc anters and cancer metastases of haematopoietic antigen genomic concers from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention of the present invention.
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Best Local S
Matches 49
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                                                                                                                                                                                                                                                                                                                                                                                                          Sequence 272 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis
                                 Human; immune; haematopoietic; immune/haematopoietic antigen; cancer, cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               AAK54951 to AAK64702 encode the human immune/haematopoietic antigen amino acid sequences given in AAM82170 to AAM91921. (I) have cytosta
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   Homo sapiens.
                                                                                 Human immune/haematopoietic antigen genomic sequence SEQ ID NO:41549
                                                                                                                      07-NOV-2001
                                                                                                                                                     AAK86737;
                                                                                                                                                                                     AAK86737 standard; DNA; 281 BP.
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                                                                                                                                                                                                                                                                                                                                                       Local Similarity
                                                                                                                                                                                                                                                                                            3073 AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT
                                                                                                                                                                                                                                                                         64
                                                                                                                                                                                                                                                                                                                                         49;
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2000US-024925P.
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2000US-0251868P.
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                                                                                                                    (first entry)
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J. 4.5e-12;
O;
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2000US-0228924P.
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polypeptides, and metastasis.

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                  GENOME
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24-FEB-2000;

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16-MAR-2000;

17-MAR-2000;

18-AR-2000;

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07-JUN-2000;

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11-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the proteins and polynucleotides may be used to prevent, diagnose and treat immune/haematopolistic-related diseases, especially
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               cancers and cancer metastases of haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoletic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 represent sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                         09-AUG-2001.
                                                                                                                                                                                                                                                                                                         WO200157182-A2.
                                                                                                                                                                                                                                                                                                                                                                      cytostatic;
                                                                                                                                                                                                                                                                                                                                                                                       Human; immune; haematopoietic; immune/haematopoietic antigen;
                                                                                                                                                                                                                                                                                                                                                                                                                       Human immune/haematopoietic antigen genomic sequence SEQ ID NO:21438.
                                                                                                                                                                                                                                                                                                                                                                                                                                                       06-NOV-2001 (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        AAK66626;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     AAK66626 standard; DNA; 21477
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 281 BP; 48 A; 79 C; 70 G; 84 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             AAK54951 to AAK64702 encode the human immune/haematopoietic antigen
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Disclosure; SEQ ID NO 41549; 3071pp + Sequence Listing; English
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                                                                                                                                                                                                                                           17-JAN-2001;
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2000US-018464P

2000US-0189874P

2000US-0199076P

2000US-0199123P

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2000US-0209467P

2000US-0214886P

2000US-0215135P

2000US-0215135P

2000US-0215137487P

2000US-02151880P
                                                                                                                                                                                                                                                                                                                                                                      gene therapy; vaccine; metastasis; ds.
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                                                                                                                                                                                                                                           2001WO-US001354
                                                                                                                                                                                          2000US-0179065P.
2000US-0180628P.
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useful f
AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis an treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent,
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                                                                                                                                                                                                                                                            Disclosure;
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                                                                                                                                                                                                                                                                                                                                                                                                                                         SCI
                                                                                                                                                                                                                                                            21438; 3071pp +
                                                                                                                                                                                                                                                                                                human immune/hematopoietic diagnosing and/or treating
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                                                                                                                                                                                                                                                                                                                                                                                                   NS.
                                                                                                                                                                                                                                                            Sequence Listing; English.
                                                                                                                                                                                                                                                                                                  antigen polypeptides, cancers and metastasis.
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11-JUL-2000
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12-JUG-2000
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RESULT 68
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ID ADZ13418 standard;
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                                                                                                                                                                                                                                                                                                                                                                                                                                             The invention relates to a nucleic acid array for detecting a cancer associated (CA) nucleic acid, comprising two or more nucleic acid probes. The invention also relates to a peptide array comprising two or more isolated polypeptides encoded by a CA nucleic acid sequence, a compound that binds to a polypeptide, an isolated antibody or its fragment which binds to a polypeptide, which is prepared by immunizing a host animal with a composition comprising the polypeptide or its antigen binding fragment and collecting cells from the host expressing antibodies against the antigen or its antigen binding fragment, a composition comprising the antibody and a carrier, a method of screening for anticancer activity, a
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          diagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metastases of haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoietic antigen genomic sequences from the present invention. AAK34942 to AAK34950 and AAM82169 represent sequences used in the exemplification of the present invention
method of detecting a CA nucleic acid, a method of diagnosing cancer, a method of treating cancer and a method of inhibiting expression of a CA nucleic acid in a cell. The CA nucleic acids are useful for detecting CA nucleic acids. The antibody is useful for detecting the presence or absence of cancer cells in an individual which involves contacting cells from the individual with the antibody and detecting a complex of a CA protein from the cancer cells and the antibody, where the detection of the complex correlates with the presence of cancer cells in the individual. The composition is useful for inhibiting growth of cancer cells in an individual or for delivering a therapeutic agent to cancer cells in an individual. The invention is also useful for diagnosing cancer, for treating cancer and for inhibiting expression of a CA gene in
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Disclosure; SEQ ID NO 938; 198pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Nucleic acid array useful for detecting cancer associated nucleic acid, comprises two or more nucleic acid probes.
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Pred. No.
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Query Match

1.6%;

Score 49;

DB 12;

Length 243428;

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RESULT 69
ADF51132/c
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                                                    target. Specifically, it refers to the guanine-nucleotide exchange factor (GEF) named P-Rexl, which has also been identified as a ghosphatidylinositol(3,4,5)P3-sensitive activator of Rac (a monomeric CP phosphatidylinositol(3,4,5)P3-sensitive activator of Rac (a monomeric CP phosphatidylinositol(3,4,5)P3-sensitive activator of Rac (a monomeric CP phosphate). Accordingly, P-Rexl can be described as having Rac-GEF activity and is adapted to function downstream of activation of heterotrimeric GP proteins in neutrophils. The present invention describes this protein as CP ceducing or inhibiting inflammation, metastasis, septic shock, reducing or inhibiting inflammation, metastasis, septic shock, corrected activities including antiinflammatory, cytostatic, annibacterial, cactivities including antiinflammatory, cytostatic, antibacterial, correction or attended antiarteriosclerotic. Furthermore, the protein or its mutant, the nucleic acid or appropriate antibody may compared in a screening assay to identify a modulator of P-Rexl binding, activity or expression. This polynucleotide is the human P-Rexl genomic CP DNA sequence of the invention.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New isolated P-Rex1 protein or its derivative useful for discovering drugs capable of reducing or inhibiting inflammation, metastasis, sej shock, neurodegeneration or atherosclerosis, or for identifying P-Re
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    human; P.Rex1; Rac; guanine-nucleotide exchange factor; GBF; GTI inflammation; metastasis; septic shock; neurodegeneration; atherosclerosis; antiinflammatory; cytostatic; antibacterial; immunosuppressive; neuroprotective; antiarteriosclerotic; gene;
Sequence 243428 BP; 65880 A; 63219 C; 59010 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 This invention relates to a novel protein useful as an anti-inflammatory
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Disclosure; SEQ
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Stephens L,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         ADF51132
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  WPI; 2004-011515/01.
P-PSDB; ADF51119.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         21-MAR-2002; 2002GB-00006684
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       21-MAR-2003; 2003WO-GB001238.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Human P-Rex1 genomic DNA sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 12-FEB-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              (BABR-) BABRAHAM INST
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Local
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49; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      85920 BP; 23268 A;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               IJ
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hes 0;
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   55319 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Indels
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              septic
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Db 235339 GATTGTTGCCACTGCACTGCACACAACAACAACAACAACACTCTGTCC 235291 RESULT 70 AAMS2458; XC 77-NOV-2001 (first entry) XC HAMARI immune/haematopoietic antigen genomic sequence SEQ ID NO:37270. XW HAMARI immune/haematopoietic antigen genomic sequence SEQ ID NO:37270. XW HAMARI immune/haematopoietic antigen; cancer; XW HOPO 9-ANG-2001. XX WO200157183-A2. XX WO200157183-A2. XX WO200157183-A2. XX W 17-AAM-2001; 20010-01790554. XX W 17-AAM-2001; 20010-01806528. PR 24-PER-2000; 200005-0186350P. PR 13-PAR-2000; 200005-0186350P. PR 13-PAR-20	Best Local Similarity 100.0%; Pred. No. 3.6e-12; Matches 49; Conservative 0; Mismatches 0; Indels Matches 49; Conservative 0; Mismatches 0; Indels 3074 GATTGTGCCACTGCACTGCAGCCTGGGCAACAGAGCAAGACTCTGTCTG
PR 08-SEP-2000 PR 11-SEP-2000 PR 21-SEP-2000 PR 21-	PR 08-SEP-2000 PR 08-SEP-2000 PR 08-SEP-2000 PR 08-SEP-2000 PR 08-SEP-2000
72000US-0232080P-0232080P-02323080P-02323080P-02323080P-0232309P-02323080P-02323080P-02323080P-02323080P-02323080P-02323080P-02323080P-02323080P-02323083P-02300US-0233083P-02323083P-02323083P-02323083P-02323083P-02323083P-02323083P-02323083P-02323083P-02323083P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-02323093P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023200US-023481P-023232P-0232321P-02323221P-02323221P-02323221P-02323221P-02323221P-02323221P-023232221P-02323222222222222222222222222222222222	2000US-0231242 2000US-0231243 2000US-0231244 2000US-0231413 2000US-0231413

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RESULT 71

AAK82461/c

ID AAK82461 standard; D

XX

AC AAK82461;

XX

DT 07-NOV-2001 (first

XX

DB Human immune/haematc

XX

KW Human; immune; haema

XX

KW cytostatic; gene the

XX
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    ARK54951 to ARK64702 encode the human immune/haematopoietic antigen (I) CC amino acid sequences given in ARM62170 to ARM91921. (I) have cytostatic CC activity, and can be used in gene therapy and vaccine production. (I) CC proteins and polynucleotides may be used in the prevention, diagnosis and CC treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat discrete associated with decreased CC expression by rectifying mutations or deletions in a patient's genome CC that affect the activity of (I) by expressing inactive proteins or to CC supplement the patients own production of (I). Additionally, (I) CC polynucleotides may be used to produce the secreted (I), by inserting the CC nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, CC diagnose and treat immune/haematopoietic-derived cells. ARK64703 CC cancers and cancer metastases of haematopoietic antigen genomic CC sequences from the present invention. ARK54921 to ARK8950 and ARM82169 CC cancers are supplied in the accentification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                    Query Match
Best Local S
Matches 48
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17.NOV-2000;
17.NOV-2000;
17.NOV-2000;
17.NOV-2000;
17.NOV-2000;
01.DEC-2000;
01.DEC-2000;
05.DEC-2000;
05.DEC-2000;
05.DEC-2000;
06.DEC-2000;
06.DEC-2000;
08.DEC-2000;
                      Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 4316 BP; 1270 A; 791 C; 832 G; 1423 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Nucleic acids encoding useful for preventing,
                                                                               Human immune/haematopoietic antigen genomic sequence SEQ ID NO:37273.
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                                                                                                                                                                                                                                                                                                                                                                                    48;
                                                                                                                                                                                                                                                                                                                                                                                                       Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       sequences used in the exemplification of the present invention
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2000US-0249265P,
2000US-024925P,
2000US-024929P,
2000US-024929P,
2000US-0250160P,
2000US-0250160P,
2000US-0251030P,
2000US-0251988P,
2000US-025173P,
2000US-0251866P,
2000US-0251866P,
2000US-0251866P,
2000US-0251869P,
2000US-0251869P,
2000US-0251989P,
2000US-0251999P,
2000US-0251999P,
2000US-025199P,
2000US-025190PP,
2000US-025190PP,
2000US-0259678P.
                                                                                                                                                                                                                                                                                                                                                                                    ilarity 100.0%;
Conservative (
                                                                                                                      (first
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                                                                                                                    entry)
                                                                                                                                                                                                                                                                                                                                                                                                                            1.5%;
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                                                                                                                                                                                                                                                                                                                                                                                    Score 48; DB;
; Pred. No. 1.20
0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                    0;
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                                                                                                                                                                                                                                                                                                                                                                                                                            DB 4;
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                                                                                                                                                                                                                                                                                                                                                                                    Gaps
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      30-JUN-2000

07-JUL-2000

11-JUL-2000

11-JUL-2000

12-JUL-2000

12-JUL-2000

14-JUL-2000

14-JUL-2000

14-JUG-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

10-SEP-2000

11-SEP-2000

21-SEP-2000

21-SEP-2000
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04-FEB-2000
24-FEB-2000
02-MAR-2000
16-MAR-2000
17-MAR-2000
17-MAR-2000
19-MAY-2000
07-JUN-2000
07-JUN-2000
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  2000US-0186628P.
2000US-0186350P.
2000US-0186350P.
2000US-0198123P.
2000US-0198123P.
2000US-0215135P.
2000US-021518867P.
2000US-0216880P.
2000US-02174867P.
2000US-0224518P.
2000US-0225513P.
2000US-0225513P.
2000US-0225513P.
2000US-0225513P.
2000US-0225513P.
2000US-0225513P.
2000US-0225513P.
2000US-022575P.
2000US-0225758P.
2000US-0225758P.
2000US-0225758P.
2000US-0225759P.
2000US-022575P.
2000US-023575P.
2000US
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2000US-0236370P. 2000US-0236802P. 2000US-0237037P. 2000US-0237038P. 2000US-0237039P.

2000US-0235834P. 2000US-0235836P. 2000US-023637P. 2000US-0236367P. 2000US-0236368P. 2000US-0236369P.

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RESULT 72
AAK82456/c
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Query Match
Best Local S
Matches 48
31-JAN-2000;
04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAX-2000;
07-JUN-2000;
28-JUN-2000;
07-JUL-2000;
07-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 4316 BP; 1270 A; 791 C; 832 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Disclosure; SEQ ID NO 37273; 3071pp + Sequence Listing; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Nucleic acids encoding
                                                                                                                                                                                                                                                                                                                                                                                                        07-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                     AAK82456;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AAK82456
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Rosen
                                                                                                                                                                                                                                          09-AUG-2001.
                                                                                                                                                                                                                                                                                                                                                                          Human immune/haematopoietic antigen genomic sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         represent sequences used in the exemplification of the
                                                                                                                                                                                                           17-JAN-2001;
                                                                                                                                                                                                                                                                        WO200157182-A2
                                                                                                                                                                                                                                                                                                                               cytostatic;
                                                                                                                                                                                                                                                                                                                                               Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         2001-483426/52
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ξ
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        preventing,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Barash SC,
                                                                                                                                                                                                                                                                                                                               gene
2000US-0179065P.
2000US-0184664P.
2000US-0184350P.
2000US-0189874P.
2000US-0198123P.
2000US-0198123P.
2000US-0205467P.
2000US-0215135P.
2000US-0215135P.
2000US-0215136P.
2000US-0215136P.
2000US-0216880P.
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llarity 100.0%; P
Conservative 0;
                                                                                                                                                                                                              2001WO-US001354
                                                                                                                                                                                                                                                                                                                                                                                                        (first entry)
                                                                                                                                                                                                                                                                                                                                 therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                 DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        human immune/hematopoietic antigen polypeptides, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Ruben
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Score 48;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  SM;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  DB
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             1423 T; 0 U; 0 Other;
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20-CCT-2000
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20-NOV-2000
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21-NOV

2000US-0237040P.
2000US-023993P.
2000US-023993P.
2000US-0240960P.
2000US-0241785P.
2000US-0241786P.
2000US-0246178P.
2000US-0246475P.
2000US-0246475P.
2000US-0246475P.
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2000US-0246528P.
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2000US-025997P.
2000US-0251989P.
2000US-025997P.
2000US-025997P.

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201-NOV-2000
08-NOV-2000
17-NOV-2000
17-NO
AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis are treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the
                                                                                                                                                                                                                         Disclosure;
                                                                                                                                                                                                                                                                                                                                                                                             (HUMA-)
                                                                                                                                                                                                                                                                                                                     2001-483426/52
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                                                                                                                                                                                                                                                              c acids encoding for preventing,
                                                                                                                                                                                                                                                                                                                                                                                              HUMAN
                                                                                                                                                                                                                                                                                                                                                          Barash
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2000US-0246476P
2000US-0246478P
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2000US-0246525P
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2000US-0246526P
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2000US-0246526P
2000US-0246519P
2000US-0246611P
2000US-0249210P
2000US-0249211P
2000US-024921P
2000US-0249211P
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2000US-0251999P
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                                                                                                                                                                                                                                                                                                                                                          sc,
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                                                                                                                                                                                                                                                                                                                                                                                              SCI
                                                                                                                                                                                                                         37268; 3071pp +
                                                                                                                                                                                                                                                          human immune/hematopoietic diagnosing and/or treating
                                                                                                                                                                                                                                                                                                                                                           MS
                                                                                                                                                                   immune/haematopoietic antigen (I) to AAM91921. (I) have cytostatic
                                                                                                                                                                                                                           Sequence
                                                                                                                                                                                                                           Listing;
                                                                                                                                                                                                                                                                cancers
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English

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polypeptides, and metastasis.

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2000US-0217487P.
2000US-0217496P.
2000US-021829P.
2000US-0224518P.
2000US-0224518P.
2000US-0225211P.
2000US-0225214P.
2000US-0225214P.
2000US-0225214P.
2000US-0225268P.
2000US-022575P.
2000US-02257182P.
2000US-023943P.
2000US-023943P.
2000US-0231243P.

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CC acid sequences from mouse and human (ADA)1482-ADA)3094), and to CC recombinant carcinoma associated proteins (CAP) encoded by them. The CC invention also encompasses expression vectors and host cells comprising a CC Anucleic acid, a polypeptide (especially an antibody) that specifically binds to the protein, and a biochip comprising CA nucleic acid or CC fragments thereof. The sequences of the invention were identified using CC oncogenic retroviruses, which insert into the genome of the host organism CC at random. Many of these do not carry transduced host oncogenes or pathogenic trans-acting viral genes, meaning that cancer incidence is a CC direct consequence of the effects of proviral integration into host protooncogenes. The CA nucleic acid sequences can be used to diagnose CC carcinoma (especially breast cancer, prostate cancer, lymphoma or leakaemia) or a propensity to carcinoma by determination of the sequence of a CA gene, or by determination of CA gene expression in particular CC tissues. CA nucleic acids, proteins and antibodies are also useful as therapeutic agents and in screening and evaluating drug candidates. The C present sequence represents a specifically claimed human CA nucleic acid sequence of the invention. Note: The complete sequence data for this complete sequence data for this sequence of the invention. Note: The complete sequence data for this complete sequence data for this cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Matches
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Best Local :
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Sequence 4317 BP; 1270 A; 791 C; 831 G; 1425 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 17-JUL-2003.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         The invention relates to recombinant carcinoma associated (CA) nucleic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 1; SEQ ID NO 1184; 245pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          New recombinant nucleic acid encoding carcinoma associated protein, useful for preparing compositions for treating carcinomas.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       WPI; 2003-587068/55
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (SAGR-) SAGRES DISCOVERY
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        26-DEC-2001; 2001US-00035832
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     26-DEC-2002; 2002WO-US041414.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human MDM2 carcinoma associated gene,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    (first
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    entry
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RESULT 75 ADE95914

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                                                                                                       Matches
                                                                                                                          Query Match
Best Local Similarity
                                                                                                                                                                                                                                                           The invention relates to a novel recombinant nucleic acid comprising a nucleotide sequence selected from any of the 660 sequences fully defined in the specification. A polynucleotide of the invention has cytostatic activity, and may have a use in gene therapy, or in a vaccine. The recombinant nucleic acids and polypeptides are useful for treating carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and may have a rescuence represents a human gene of the invention.
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23-CCT-2001; 2001US-00004113.
08-NOV-2001; 2001US-00052482.
30-NOV-2001; 2001US-00997722.
20-DEC-2001; 2001US-00034650.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       human; ds; cytostatic; gene therapy; vaccine; carcinoma; lymphomas; cancer; neoplasm; adenocarcinoma; sarcoma; gene.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human MDM2 gene
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            ADB72404;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            ftp.wipo.int/pub/published_pct_sequences.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    30-JAN-2003
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                                                                                                                                                                                                       Sequence 52242 BP; 14384 A; 10354 C; 10997 G; 16487 T; 0 U;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Claim 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              WPI; 2003-239337/23
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New recombinant nucleic acid, useful for treating carcinomas, lymphomas
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             26-DEC-2001; 2001WO-US051291
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                                GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG
                                                                                                                                                                                                                                                                                                                                                                                                                                                                        SEQ ID NO 232; 2304pp; English.
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                                                                                                       Conservative
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                                                                                                                                Score 48; DB 10;
Pred. No. 1.1e-1;
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Pred. No.
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                                                                                                                                   1.1e-11;
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RESULT 76
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Best Local Similarity
Matches 48; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         diagnosis and treatment of cancer, especially carcinomas, as well as the use of compositions in screening methods. The compositions of the invention may have cytostatic activity whilst the disclosed sequences may be useful for gene therapy. The carcinoma associated nucleic acids and proteins are useful for diagnosing and treating carcinomas, for example lymphoma, breast cancer, prostate cancer or leukaemia, or for screening drug candidates or bioactive agents capable of binding to, or modulating the activity of, a carcinoma associated protein. The present sequence is the genomic DNA sequence of the human MDM2 gene which is a carcinoma
                                                                                                                                                                                                                                          Human ENTPD5 gene genomic sequence SEQ ID NO:85.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Sequence 52242 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Claim 1; SEQ ID NO 172; 793pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    New carcinoma associated nucleic acids and proteins, useful for screening drug candidates, or for diagnosing and treating carcinomas, e.g. lymphoma, breast cancer, prostate cancer or leukemia.
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lymphoma; breast
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      16-JUN-2005
                                                                                                                       Homo sapiens
                                                                                                                                                                               DNA methylation;
                                                                                                                                                                                                                                                                                                      25-AUG-2005
                                                                                                                                                                                                                                                                                                                                                                                                                             AEA61175 standard; DNA; 53779 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       associated gene of the invention.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    This invention relates to novel recombinant nucleic acids for use
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (SAGR-) SAGRES DISCOVERY.
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Conservative
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                                                                                                                                                                               biomarker; cancer; gene; ds; ENTPD5
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           14384 A; 10353 C; 10998 G; 16487 T; 0 U; 20 Other;
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Pred. No.
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1.1e-11;
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RESULT 77
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   CC nucleic acid sequences useful as a biomarker for a disease to be competited acid sequences useful as a biomarker for a disease to be comprising comethylated (M1) involves identifying nucleic acid sequences comprising comethylated in diseased cells, comparing expression level of nucleic acid sequences with that of demethylated nucleic acid sequences and comparing expression level of nucleic acid sequences exhibiting increase in expression comparing comparing expression level of nucleic acid sequences and comparing expression of a disease in a subject, which involves determining the degree of methylation of one or more CpG sites on nucleic acid sequences in a composition to, or stage of the disease in the subject based on the degree of methylation; (2) monitoring the onset, progression, or compound for inhibiting a disease in the subject based on composition to or expression of a disease in a subject; (3) determining the efficacy of a composition of a disease in a subject; and (4) a kit (1) composition of a disease in a subject; and therapeutic compound for inhibiting a disease in a subject; and (4) a kit (1) composition of a disease in a subject; and therapeutic treatment of a disease. (M1) is useful for identifying one or more composition to a subject; and therapeutic composition of a disease useful for identifying one or more composition. The present sequence sequences are useful for detecting, the present sequence represents a specifically claimed human composition. The present sequence represents a specifically claimed human control of the invention. Note - The sequence data for this patent is not represented in the printed control of the composition but was obtained in electronic format from the USPTO web
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Query Match
Best Local Similarity
Matches 48; Conserv
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                                                                                                                                                                                                                                                                            Human genomic sequence hCG16651.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Sequence 53779 BP; 14286 A; 11767 C; 12248 G; 15478 T; 0 U; 0 Other;
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                                                                     WO2003073826-A2
                                                                                                                                        Homo sapiens.
                                                                                                                                                                                                              Cytostatic; carcinoma; lymphoma;
                                                                                                                                                                                                                                                                                                                                                      18-NOV-2004
                                                                                                                                                                                                                                                                                                                                                                                                                            ACN44374;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          ACN44374 standard; DNA; 181684 BP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             demethylated nucleic
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            16-DEC-2003; 2003US-00737082
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1.5%; So illarity 100.0%; I Conservative 0;
                                                                                                                                                                                                                                                                                                                                                  (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Score 48;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Mismatches
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                                                                                                                                                                                                              cancer;
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. 1.1e-11;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Length 53779;
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Best Local
                                                                                       16-MAR-2000;
25-MAY-2000;
09-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bloactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bloactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing carcinoma, (vi) for inhibiting the activity of CAP; (ii) for tracting carcinoma; (viii) for neutralizing the effect of CAP; (vii) for tracting; (x) for diagnosing carcinoma or a propensity to carcinoma, and (xi) for determining Carcinoma accinoma or a propensity to carcinoma. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              WPI; 2003-328604/31
                                                                                                                                    17-FEB-2000;
                                                                                                                                                                                                                                                                                                                                        Human prostate expression marker cDNA 16322.
                                                                                                                                                                                                                                                                                                                                                                       13-SEP-2002
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Claim 1; SEQ ID NO 790; Opp; English.
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                              (MILL-)
                                                                                                                                                                  20-FBB-2001; 2001WO-US005171.
                                                                                                                                                                                                                                 WO200160860-A2
                                                                                                                                                                                                                                                                                          Human; prostate cancer; cytostatic; carcinogen; pharmacodyanamic marker; pharmacogenomic marker; gene; ss.
                                                                                                                                                                                                                                                                                                                                                                                                       ABV16331;
                                                                                                                                                                                                                                                                                                                                                                                                                                  ABV16331 standard; cDNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                sequence. Note: This patent is an equivalent to basic patent
US2002182586A1, for which no sequence data was published
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                              MILLENNIUM PREDICTIVE MEDICINE INC.
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                                                         ; 2000US-0183319P.
; 2000US-0189862P.
; 2000US-0207454P.
; 2000US-0211314P.
; 2000US-0219007P.
; 2000US-0255281P.
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                                                                                                                                                                                                                                                                                                                                                                       (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      1.5%; Score 48; DB
100.0%; Pred. No. 1.:
tive 0; Mismatches
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Monahan JE
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1.1e-11;
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RESULT 79
ABV46129
ID ABV46
XX ABV46
XX ABV46
XX IN-SE
DT 16-SE
XX Human
XX Pharm
XX Pharm
XX Pharm
XX Pharm
XX Pharm
XX IN-FE
PR 15-MA
PR 25-MA
PR 18-JU
PR 11-EE
PR 11-U
PR 
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                                                                                                                                                                                                                                                                                                                                                                                                                                            16-MAR-2000;
25-MAY-2000;
09-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        a patient is afflicted with prostate cancer; (b) monitoring the progression of prostate cancer in a patient; (c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient; (d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient; (e) selecting a composition for inhibiting prostate cancer in a patient; (f) assessing the prostate cell carcinogenic potential of a compound; (g) determining whether prostate cancer has metastasized in a patient; (h) assessing the aggressiveness or indolence of prostate cancer in a patient; (l) is also useful as a pharmacodyanamic or pharmacogenomic marker
                                                                                        Novel isolated nucleic acid molecule associated with cancerous state of prostate cells and correlating with presence of prostate cancer, useful for detecting presence of prostate cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human; prostate pharmacogenomic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     17-FEB-2000;
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                                                                                                                                                                                                                                                                                                                              (MILL-) MILLENNIUM PREDICTIVE MEDICINE
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                                                                                     detecting
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               3070 GCAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 3116
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       prostate expression marker cDNA 46120.
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                                                                                                                                                                                                                                                                                                                                                                                  ; 2000US-0183319P.
; 2000US-0189862P.
; 2000US-0207454P.
; 2000US-0211314P.
; 2000US-0219007P.
; 2000US-0255281P.
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                                                                                     presence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 cancer; cytostatic; carcinogen; pharmacodyanamic marker;
marker; gene; ss.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               1.5%;
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                                                                                                                                                                                                                                                                       Monahan JE
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Score 47;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        ВÞ
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  DB 5; Le
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9110; 11750pp; English

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RESULT 80
ADL13941
ID ADL13941
XX ADL133
AC ADL133
AC OSTEO
XX OST
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The invention relates to a method of determining susceptibility of an individual to joint space narrowing and/or seteophyte development and/or joint pain comprising identifying whether the individual has at least one polymorphism in a polymucleotide encoding at least one of the protein listed in the specification. The methods, composition and agent are useful for modulating the susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain that is associated with a disease, preferably osteoarthritis. The cell line and the non-human animal are useful for screening for an agent for diagnosing an individual having susceptibility to joint space narrowing and/or osteophyte development and/or joint space narrowing and/or osteophyte development and/or joint space narrowing and/or to the polynucleotide encoding a protein listed in the specification. (Note:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Disclosure; SEQ ID NO 473; 297pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Jones
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joint space narrowing; osteophyte development; joint pain;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Osteoarthritis-associated polymorphic nucleotide
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Pred. No.
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Best Local
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The invention comprises novel human gene trapped sequences (GTSs) which are useful in gene discovery and as markers for gene expression analysis, forensic analysis, and determining the genetic basis of human disease. The human GTSs of the invention are useful for diagnosing and treating disorders affecting development and cell differentiation, such as: aging, cancer, autoimmune disease, hupus, scleroderma, Crohn's disease, multiple sclerosis, inflammatory bowel disease, immune disorders, schizophrenia, psychosis, alopecia, glandular disorders, inflammatory disorders, ataxia telangiectasia, diabetes, skin disorders, osteoarthritis, rheumatoid
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    The sequence data for this patent did not form specification but was obtained in electronic for ftp.wipo.int/pub/published_pct_sequences).
                                                                                                                                                                                                           New human gene trapped sequences, useful for diagnosing and treating disorders affecting development and cell differentiation, e.g. aging, cancer, schizophrenia, alopecia, diabetes, rheumatoid arthritis, or infertility.
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                                                                                                                                                                          Claim 3;
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27-OCT-1999;
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47; Conserv
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Pred. No.
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and immunostitudiates, and a polypeptide (III) having a CDNA sequence (S1) of a ovarian tumour collection the 10912 nucleotide sequences as given in ABL77023 to ABL87934, (III) encoding (II) having a sequence (S2), a T cell population of (II), or antigen presenting cells that express (II). (I) has cytostatic activity. An oligonucleotide (IV) that hybridises to (S1) can be used for detecting ovarian cancer in a patient's biological sample preferably serum or ovarian tissue. The method comprises contacting a biological sample from a patient with (IV), detecting the amount of polynucleotide hybridising to (IV) and comparing the amount to a predetermined cutoff value and thereby detecting ovarian cancer in the patient, where the amount of polynucleotide hybridising to (IV) is detected preferably by collecting comparison of rise detected preferably by collecting and/or expanding T cells specific for an ovarian tumour protein comprising contacting T cells with (III) or (II) is useful in design and preparation of ribozyme molecules for inhibiting expression of the tumour polypeptides and proteins in tumour cells; and to isolate a full length gene from a suitable library e.g., a tumour cDNA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Query Match
Best Local &
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       polypeptide of a ovari polypeptide, antibody polypeptide.
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46; Conserv
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25-MAY-2000;
15-JUN-2000;
                                patient afflicted with ovarian cancer comprising providing to cells of the patient an antisense oligonucleotide complementary to a marker of the invention. The markers are useful for assessing if a patient is afflicted with ovarian cancer, which involves comparing the level of expression of a marker in a patient sample and a normal level of expression of a marker in a control non-ovarian cancer sample. A difference between the expression levels indicates ovarian cancer. The level of expression of a marker corresponds to a secreted protein or to a transcribed polynucleotide or its portion. The level of expression of the marker is assessed by detecting the presence in the sample, a protein or protein fragment is detected using an antibody that specifically binds with the protein or p
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   The invention relates to nucleic acid markers which are overexpressed in ovarian cancer cells as compared to their expression in normal (i.e. non-cancerous) ovarian cells. The invention also relates to polypeptides encoded by the markers, antibodies that selectively bind to the polypeptides, a method of inhibiting ovarian cancer in a patient at risk of developing ovarian cancer involving inhibiting expression of a gene corresponding to a marker of the invention and a method of treating a
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Novel isolated nucleic acid molecules (markers) overexpressed in ovarian cancer cells as compared to their normal non-cancerous ovarian cells are used to characterize stage, grade, histological type of ovarian cancer.
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25-JUL-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           library using well known techniques
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Disclosure; SEQ ID NO 17260; 106pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     WO200170979-A2
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGACACAGACTC 3116
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2000US-0211940P.
2000US-0216820P.
2000US-0220661P.
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Pred. No.
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RESULT 84
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Best Local (
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25-MAY-2000;
09-JUN-2000;
a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the specification or its complement. (I) is useful for: (a) assessing whether a patient is affilicted with prostate cancer; (b) monitoring the progression of prostate cancer in a patient; (c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient; (d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient; (e) selecting a composition for inhibiting prostate cancer in a patient; (f) assessing the prostate cancer has metastazed in a patient; (h) determining whether prostate cancer has metastazed in a patient; (h) assessing the aggressiveness or indolence of prostate cancer in a patient
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         polynucleotide which anneals with the marker or anneals with a portion of the polynucleotide comprising the marker, under stringent conditions. The marker is also used for monitoring the progression of ovarian cancer in a patient which involves detecting expression of the marker in a patient sample at a first point in time, repeating the method at a subsequent time and comparing the level of expression. The method is carried out using an ovarian tissue sample. A composition comprising a marker, polypeptide or antibody of the invention is used to treat ovarian cancer. This sequence represents a human ovarian cancer DNA marker of the
                                                                                                                                                                                                                                                   The invention relates to
                                                                                                                                                                                                                                                                                                    Claim 1;
                                                                                                                                                                                                                                                                                                                                                                          Novel isolated nucleic acid molecule associated with cancerous state prostate cells and correlating with presence of prostate cancer, usef
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                                                                                                                                                                                                                                                                                                 Page 11527; 11750pp; English.
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; 2000US-0189862P.
; 2000US-0207454P.
; 2000US-0211314P.
; 2000US-0219007P.
; 2000US-0255281P.
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RESULT 85
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11-JAN-2000;
02-MAY-2000;
           oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide which comprises a 1-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH13642 represent human cDNA sequences; AAB92446 to AAB95893 represent human and AAH13630 to AAH13629 represent human cDNA sequences; AAB92446 to AAB95893
                                                                                                                                                                                                                                                                                                                                                                                                The present invention describes primer sets for synthesising 5602 full-length CDNAs defined in the speciation. Where a primer set comprises:

(a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Primer sets for synthesizing polynucleotides, particularly the 5602 length cDNAs defined in the specification, and for the detection and diagnosis of the abnormality of the proteins encoded by the full-length control of the proteins encoded by the protein
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Claim
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46; Conser
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2000JP-00118776.
2000JP-00183767.
2000JP-00241899.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       NO 18263;
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, 100.0%; Pr
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T, Wakamatsu
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Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Sequence
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C, Otsuki
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RESULT 86
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RESULT 87
ACC82887/c
ID ACC82887 standard; DNA; 7001 BP
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Matches 46
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Best Local
                                                                                                                                                                                                                                                          The invention relates to an isolated nucleic acid encoding a PRO polypeptide. The polypeptide, agonist or an antagonist, antibody, composition, and method are useful for diagnosing and treating an related disorder, e.g. systemic lupus crythematosus, rheumatoid arthritis. The present sequence represents a DNA encoding a PRO
                                                                                                                                                                                                                                                                                                                                                                                           New nucleic acid encoding PRO polypeptide, useful for diagnosing and treating an immune related disorder, e.g. systemic lupus erythematosus, rheumatoid arthritis, osteoarthritis, thyroiditis, or diabetes mellitus
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                                                                                                                                                                                                                                                                                                                                                            Claim 1; SEQ ID NO 1453; 158pp; English.
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                                                                                                                                                                                                              Sequence 6530 BP; 1644 A; 1400 C; 1429 G;
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                                                                                                                                               1 Similarity
46; Conser
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                                                                                 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 3544
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Clark H,
                                                                                                                                               1.5%;
ilarity 100.0%;
Conservative
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b; Pred. No. 1e-
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Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human thyroid hormone receptor interactor 6 (TRIP6) gene
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                  05-NOV-2002; 2002WO-US035479
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3708. .3877
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Best Local :
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16-MAR-2000;
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07-JUN-2000;
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11-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        The invention relates to antisense compounds targetted to a nucleic acid encoding thyroid hormone receptor interactor 6 (TRIP6) to inhibit its expression. TRIP6 is also known as OPA-interacting protein-1 (OIP-1) and zyxin-related protein-1 (ZRP-1). TRIP6 DNA is located on chromosome 7q22. Antisense compounds of the invention are useful for modulating the expression of TRIP6 and for treating diseases or conditions associated with the expression of TRIP6 such as hyperproliferative disorders (e.g. cancer). They are useful for diagnostics, therapeutics, prophylaxis e.g. to prevent or delay infection, inflammation or tumour formation, as research research research sand kits and in distinguishing between functions of various members of a biological pathway. The are also useful in antisense therapy. The present sequence is human TRIP6 gene fragment
                                                                                                                                                                                                                                                                                        02-AUG-2001.
                                                                                                                                                                                                                                                                                                                         WO200155320-A2
                                                                                                                                                                                                                                                                                                                                                          Homo
                                                                                                                                                                                                                                                                                                                                                                                         cancer;
                                                                                                                                                                                                                                                                                                                                                                                                       Human; reproductive system related antigen; reproductive system disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                        Human reproductive system related antigen DNA SEQ ID NO:
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2000US-0179065P.
2000US-0184664P.
2000US-0186350P.
2000US-0186350P.
2000US-0199076P.
2000US-0199076P.
2000US-0298132P.
2000US-0215815P.
2000US-0214886P.
2000US-021647P.
2000US-021647P.
2000US-021647P.
2000US-0217487P.
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21-SEP-2000;
21-SEP-2000;
25-SEP-2000;
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27-SEP-2000;
27-SEP-2000;
29-SEP-2000;
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22-AUG-2000;
22-AUG-2000;
23-AUG-2000;
30-AUG-2000;
01-SEP-2000;
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05-SEP-2000;
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06-SEP-2000;
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08-SEP-2000;
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14-SEP-2000;
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   2000US-0218290P

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2000US-0224819P

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                                                                                                                                                                                     The present invention provides the protein and coding sequences of a number of human reproductive system related antigens. These can be used in the prevention and treatment of reproductive system disorders, including cancer. The present sequence is a genomic sequence encoding a protein of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                               Isolated nucleic acid molecule encoding a reproductive system antigen used in preventing, treating or ameliorating a medical condition.
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                                                                                                                                Sequence
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2000US-0246611P.
2000US-0249213P.
2000US-0249211P.
2000US-0249213P.
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2000US-02511869P.
2000US-0251198P.
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live 0; Mismatches
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2000US-0179065P

2000US-0184664P

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2000US-0184664P

2000US-0199076P

2000US-0199076P

2000US-0214866P

2000US-0216647P

2000US-0216880P

2000US-02174967P

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2000US-022573PP

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Best Local Similarity
Matches 46; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New isolated ovarian polypeptides, useful
Human; connective tissue related disorder; cytostatic; gene; ds.
                                                 Genomic
                                                                                 21-MAY-2002
                                                                                                                                               ABK43029
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Disclosure; SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        WPI; 2001-488786/53.
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                                                 sequence #928 encoding novel human connective tissue polypeptide.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    ated ovarian and/or breast cancer related nucleic acids and ides, useful for diagnosing, treating and/or preventing humand disorders, particularly ovarian and/or breast cancer.
                                                                                                                                                                                                                               CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 12166
                                                                                                                                              standard; DNA; 18501
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                                                                                                                                                                                                                                                                                                           Score 46;
Pred. No.
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                  cancer; gene therapy;
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2000US-024924.P.

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Best Local
                                                                                                                                                                                                          atherosclerosis; myocarditis; cardiopulmonary bypass complication; autoimmune disease; multiple sclerosis; allergic reaction; asthma; rhinitis; eczema; inflammatory condition; Crohn's disease; nephritis; gastrointestinal disorder; inflammatory bowel disease; organ transplant rejection; immune system disorder; Bruton's disease; x-linked lymphoproliferative disorder; HIV; AIDS; infection; becall lymphoproliferative disorder; HIV; AIDS; infection; chromosome identification; chromosome mapping;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           The present invention relates to the isolation of novel human connective tissue related polypeptides (ARU86435-ARU86923) and the polynucleotide (CDNA and genomic) sequences encoding them. The sequences of the invention are useful in the diagnosis, treatment, prevention and/or prognosis of diseases associated with connective tissue(s), including cancer. The polynucleotide sequences of the invention are also useful in gene therapy. ABK42102-ABK43116 represent genomic sequences encoding the novel human connective tissue related polypeptides. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                          antiarteriosclerotic; immunosuppressive; antirheumatic; antiarthritic; antiarteriosclerotic; immunosuppressive; antirheumatic; dermatological; antiastematic; dermatological; nephrotopic; virucide; funglcide; antibacterial; antiparasitic; gene therapy; ds; connective tissues disorder; rheumatoid arthritis; systemic lupus erythematosus; scleroderma; Sjogren's syndrome; cancer; cancer metastasis; neoplasia; leukaemia; neurodegenerative disorder; hlzheimer's disease; Parkinson's disease; cardiovascular disease; hlzheimer's disease; parkinson's disease; cardiovascular disease;
 31-JAN-2000;
04-FEB-2000;
                                                      07-MAR-2002; 2002US-00092154.
                                                                                                                                                               Homo sapiens.
                                                                                                                                                                                               connective tissue related polynucleotide; gene;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           cytostatic; neuroprotective; nootropic; antiparkinsonian; cardiovascular;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Connective tissue related genomic DNA #928.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Nucleic acid encoding novel connective tissue associated polypeptides, used in diagnosing, preventing, treating or ameliorating a disorder such as cancer or rheumatoid arthritis.
                                                                                                                             US2003054375-A1
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             standard; DNA; 18501 BP
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ilarity 100.0%;
Conservative
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 2000US-0179065P.
2000US-0180628P.
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J. 9.9e-11;
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02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
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07-JUN-2000;
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2000US-0209467P.
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                   New
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connective tissue-related polypeptides and polynucleotides, useful treating, preventing and/or prognosing e.g. disorders of connective sue, (e.g. rheumatoid arthritis), cancers, cancer metastases and/or
                                                                    2003-634869/60
DB; ADB59732.
                                                                                                                                                       HUMAN
                                                                                                                      Ruben SM,
                                                                                                                                                                                                                                                      2000US-0237039P.
2000US-0237040P.
2000US-023993FP.
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2000US-024921P.
2000US-025198P.
2000US-025186P.
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                                                                                                                                                                                         2001US-00764847.
                                                                                                                                                       GENOME
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                                                                                                                       Barash
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RESULT 92
ADC210/C
ID ADC210
XX ADC210
XX ADC210
XX ADC210
XX ADC210
XX BORNA
AC ADC210
XX IN-
XX Gene t
KW Inflam
XX Infect
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P
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               S
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The invention describes an isolated nucleic acid molecule (I), which comprises a sequence that is at least 95 % identical to a connective tissue-related polynucleotide encoding connective tissue antigens (CTA). The polypeptide or polynucleotide is useful for preventing, treating, or ameliorating medical conditions in a mammal. The connective tissue polynucleotides and antibodies are particularly useful for treating, preventing and/or prognosing disorders of connective tissues (e.g. rheumatoid arthritis, discoid and systemic lupus erythematosus, cscleroderma, or Sjogren's syndrome), cancers, cancer metastases and/or neoplasias (e.g. leukaemia), neurodegenerative disorders (e.g. atherosclerosis, myocarditis or cardiopulmonary bypass (e.g. atherosclerosis, myocarditis or cardiopulmonary bypass (complications), autoimmune diseases (e.g. systemic lupus erythematosus, rheumatoid arthritis, or multiple sclerosis), allergic reactions (e.g.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Query Match
Best Local S
Matches 46
                                                                                                                                                                                                                                                                                                                                                                                                                27-MAR-2001;
12-SEP-2001;
12-SEP-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   gene therapy; human; secreted protein; haemopoietic disorder; haematological disorder; anaemia; haemophilia; inflammatory disease; cancinflammatory bowel disease; Crohn's disease; neophastic disease; cancileukaemia; wound healing; epithelial cell proliferation disorder; immune disorder; autoimmune disorder; asthmatic disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Disclosure;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      neoplasias.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ADC21019
                                                                                                                                      New human secreted proteins and nucleic acid molecules, useful for preparing a diagnostic or pharmaceutical composition for diagnosing preventing or treating hematopoietic or hematologic disorders, e.g
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        gastrointestinal
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  cardiovascular disorder; atherosclerosis; myocarditis;
infectious disease; HIV; AIDS; endocrine disorder; diabetes;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human secreted protein-related
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ADC21019;
                                                                                                                  anemia
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               26-MAR-2002;
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                                                                                                                                                                                                                                                                                                                                                                  (HUMA-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  sapiens.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           standard;
                                                                                                                  hemophilia
                                                                                                                                                                                                                                                                                                            Ruben
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ilarity 100.0%;
Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                  2001US-0278650P.
2001US-00950082.
2001US-00950083.
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                                                                                                                                                                                                                                                                                                                                                                  GENOME
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                                                                                                                                                                                                                                                                                                            SM;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           disorder;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           DNA; 18501
                                                                                                                                                                                                                                                                                                                                                                  SCI INC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        duodenal ulcer; gastroenteritis; gene;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  248pp;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Score 46; DB; Pred. No. 9.9.
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                                                                                                                                                                        diagnosing
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                                                                                                                                                 e.g.
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Disclosure;

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973; 1512pp; English

invention comprises the amino acid and coding sequences

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RESULT 93
ABT17021/c
LD ABT17021 standard; DNA; 18501
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Best Local :
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The invention comprises the amino acid and coding sequences of human secreted proteins. The DNA and protein sequences of the invention are useful for the diagnosis and treatment of allergic disorders, asthmatic disorders and immediate hypersensitivity diseases (e.g. hay fever, allergic conjunctivitis and allergic rhinitis). The proteins of the invention are also useful for identifying a binding partner. The nucleic acids of the invention are also useful for chromosome identification,
                                                                                                                                                                                                                                                                                                                                  Use of human secreted proteins and nucleic acids for preparing a diagnostic or pharmaceutical composition for diagnosing or treating allergic or asthmatic disorders, e.g. asthma, hay fever, or allergic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        27-MAR-2001; 2001US-0278650P.
12-SEP-2001; 2001US-00950082.
12-SEP-2001; 2001US-00950083.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; gene; ds; protein therapy; immediate hypersensitivity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Sequence 18501 BP; 5301 A; 3948 C; 3746
                                                                                                                                                                                                                                             Disclosure; Page 810-815; 823pp; English
                                                                                                                                                                                                                                                                                                         conjunctivitis
                                                                                                                                                                                                                                                                                                                                                                                                                                                               WPI; 2003-175010/17.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        allergic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human secreted
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        03-APR-2003
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   (HUMA-) HUMAN GENOME SCI INC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   llergic disorder; asthmatic disorder; gene therapy; secre ay fever; allergic conjunctivitis; allergic rhinitis; inding partner identification; chromosome identification;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             hybrid mapping; long-range restriction
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Ruben
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (first
                                                                                                                                                                                                                                                                                                         or rhinitis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         protein-related DNA sequence - SEQ
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           , MS
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     1.5%;
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     DB 10; ]
9.9e-11;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          mapping
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             ID No
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    disease;
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RESULT 94
ABZ68161/G
ID ABZ68
XX ABZ68
XX ABZ68
XX Y
XX Human
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                                                 The invention relates to novel human genes (ABZ66891-ABZ68209) and the CC encoded secreted proteins (ABP9470-ABP9872) useful for preventing, treating or ameliorating medical conditions e.g. by protein or gene CC treating or ameliorating medical conditions e.g. by protein or gene CC therapy. The genes are isolated from a range of human tissues disclosed in the specification. The nucleic acids, proteins, antibodies and CC (ant)agonists are useful in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast and ovarian cancer and other cancers of the CC (alcancer, e.g. breast and ovarian cancer and other cancers of the CC (alcancer, e.g. breast and ovarian cancer e.g. Addison's disease, lung or urogenital; (b) immune disorders e.g. Addison's disease, CC (allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis, CC diabetes mellitus, Crohn's disease, multiple eclerosis, rheumatoid CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as CC myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g. Carterial formal and pellepsy; and (f) infectious diseases such as viral,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Matches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New human secreted proteins encoded by genes contained in cDNA clones (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AI multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis o
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      antifungal; antiparasitic; cardiant; immune disorder; infection; vaccine; cardiovascular disorder; neurological disease; nephrotropic;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human; secreted protein; nootropic; neuroprotective;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    ABZ68161
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                West Nile fever.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            WPI; 2003-040583/03
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12-SEP-2001;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Page 2321-2325; 2423pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Conservative
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antiparkinsonian; antisickling; antis
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Pred. No.
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. 9.9e-11;
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Human

purinergic receptor P2X4 gene sequence

numan;

gene; ds.

fat deposition; leanness; non-insulin dependent diabetes mellitus; NIDDM; purinergic receptor; P2X4; antidiabetic; anorectic; diabetes; obesity;

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RESULT 96
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        New antisense oligonucleotides for modulating P2X4 expression, diagnosing, preventing or treating conditions associated with neurological disorders, osteoporosis or rheumatoid arthritis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          The invention comprises antisense oligonucleotides that are targeted to a nucleic acid encoding P2X4. The antisense oligonucleotides are useful for inhibiting the expression of P2X4 in cells or tissues to treat diseases associated with P2X4 expression, such as: neurological disorders, bone disorders (e.g. osteoporosis), or rheumatoid arthritis. The present nucleic acid represents the human P2X4 genomic DNA sequence.
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9.8e-11;
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Matches 46
                                                                                This invention relates to a novel method of diagnosing a predisposition to fat deposition, leanness or non-insulin dependent diabetes mellitus (NIDDM) in a subject. The method comprises detecting the presence or absence of a polymorphic variation associated with fat deposition, leanness or NIDDM at a polymorphic site in a purinergic receptor (P2X4) nucleotide sequence in a nucleic acid sample from a subject. The invention may be useful for the development of compounds with an antidiabetic or anorectic activity. The method is useful for diagnosing a predisposition to fat deposition, leanness or NIDDM. The nucleic acid encoding the polypeptide is useful for diagnosing conditions or diseases including fat deposition or NIDDM, also in treating diabetes and obesity. The present sequence is that of the purinergic receptor (P2X4) nucleotide sequence which was used in the method of the invention.
                                                                                                                                                                                                                                                                                                                 Diagnosing predisposition to fat deposition, leanness or non-insulin dependent diabetes mellitus (NIDDM) comprises detecting the presence absence of a polymorphic variation in a purinergic receptor.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        variation
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variation
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                                                         Sequence 28616 BP;
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                                                                                                                                                                                                                                                                                        Claim 1; SEQ ID NO 1; 154pp; English.
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  46;
                 Similarity
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  Conservative
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replace(14744,T)
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replace(22713,T)
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replace(21708,G)
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replace(15847,T)
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replace(15059,G)
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replace(17338,T)
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                Score 46; DB 12; Pred. No. 9.7e-11;
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2895 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940

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4783

GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 4738

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RESULT 97
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ID ADZ42284 standard;
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This invention relates to a novel method for testing hypertensive renal
               Claim 1; SEQ ID NO 11; 440pp; Japanese.
                                      Testing hypertensive renal disease factor, by determining polymorphism in genotype of gene relevant to hypertensive renal disease, and estimating risk factor for hypertensive renal disease based on determined genotype,
                                                                       WPI; 2005-326228/34.
                                                                                                              09-OCT-2003; 2003JP-00350959
                                                                                                                              09-OCT-2003; 2003JP-00350959
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                 disease; nephrotropic; SNE
nucleotide polymorphism;
                                                                                      KOKURITSU JUNKANKI BYO CENT SOCHO.
DOKURITSU GYOSEI HOJIN IYAKUHIN IRYO KIK.
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sm; SNP; Klotho;
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Best Local Similarity 100.0%; F
Matches 46; Conservative 0;
to a method for treating cancers involving administering to a patient an inhibitor of CAP, and a method of screening for anticancer activity in a potential drug involving providing a cell that express a CA gene, contacting a tissue sample derived from a cancer cell with an anticancer drug candidate and monitoring the effect of the anticancer drug candidate on expression of the CA gene. The CAP proteins are useful for detecting cancer associated with expression of a CAP protein in a test cell sample and for screening for a bioactive agent capable of modulating the activity of a CAP protein. The CA nucleic acids are useful for diagnosing
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       adrenoreceptor, aldosterone synthetase, endothelium nitrogen monoxide synthetase, klotho and a sodium-calcium exchanger. Furthermore, it provides primers and probes for determining hypertensive renal disease factors, in particular in relation to renal diseases including hypertensive early renal disease and hypertensive kidney blood flow obstruction. The method enables detection of risk factors, and thus hel in preventing or delaying renal disease. This polynucleotide sequence is the full length human Klotho gene given in an exemplification of the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   genotype of a gene relevant to hypertensive renal disease and estimating the risk factor for developing the disease accordingly. The present invention describes identifying gene accordingly. The present invention describes identifying the disease accordingly. The present invention describes identifying the disease in at least one of the following genes, namely endothelin converting-enzyme 1, mineralocorticoid receptor, urotensin II, superoxide-dismutase 3, thiazide sensitivity NaCl symporter, guanosine cyclase-A, hepatocyte growth factor, beta-3
                                                                                                                                                                                                  The invention relates to cancer-associated proteins (CAP) and the cancer-associated (CA) nucleic acids encoding them. The invention also relates
                                                                                                                                                                                                                                                                                                          Novel human cancer associated protein encoded within open reading of cancer associated gene, useful as targets for diagnosing cancer
                                                                                                                                                                                                                                                                                                                                                                           WPI; 2004-499109/47
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                                                                                                                                                                                                                                                                 Claim 16; SEQ ID NO 526; 182pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                    Morris DW,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     17-DEC-2002; 2002US-00322281
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human cancer-associated (CA)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                               (SAGR-) SAGRES DISCOVERY INC
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                                                                                                                                                                                                                                                                                                                                                                                                                      Malandro
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Pred. No.
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                                                                                                                                                                                                                                                                                                            cancer.
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cc associated (CA) nucleic acid, comprising two or more nucleic acid probes.

Cc The invention also relates to a peptide array comprising two or more

cc isolated polypeptides encoded by a CA nucleic acid sequence, a compound

cc that binds to a polypeptide, which is prepared by immunizing a host animal

cc with a composition comprising the polypeptide or its fragment which

cc with a composition comprising the polypeptide or its antigen binding

cc fragment and collecting cells from the host expressing antibodies against

cc the antigen or its antigen binding fragment, a composition comprising the

cc antibody and a carrier, a method of screening for anticancer activity, a

cc method of detecting a CA nucleic acid, a method of diagnosing cancer, a

cc method of treating cancer and a method of inhibiting expression of a CA

cc nucleic acids. The antibody is useful for detecting the presence or

cc absence of cancer cells in an individual which involves contacting cells

cc protein from the cancer cells and the antibody, where the detection of

cc the complex correlates with the presence of cancer cells in an individual. The composition is useful for inhibiting growth of cancer

cc cells in an individual or for delivering a therapeutic agent to cancer

cc cells in an individual. The invention is also useful for diagnosing
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Matches
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Best Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      The invention relates to a nucleic acid array for detecting a cancer associated (CA) nucleic acid, comprising two or more nucleic acid pr
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Disclosure; SEQ ID NO 60; 198pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             comprises two or more nucleic acid probes.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Nucleic acid array useful for detecting cancer associated nucleic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Diagnosis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human cancer-associated genomic DNA #7.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 58922 BP; 13257 A; 15256 C; 16255 G; 14082 T; 0 U; 72
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Morris DW,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               23-SEP-2003; 2003US-00669920
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               23-SEP-2004; 2004WO-US031617
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       cytostatic;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             ftp.wipo.int/pub/published_pct_sequences
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          DNA microarray; microarray; biochip; cancer; neoplasm;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          gene;
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cancer and
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for inhibiting expression of
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                DB 13; ; 9.5e-11;
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RESULT 101

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Best Local Similarity
                                                                                                                                                  diagnosis and treatment of cancer, especially carcinomas, as well as the use of compositions in screening methods. The compositions of the invention may have cycostatic activity whilst the disclosed sequences may be useful for gene therapy. The carcinoma associated nucleic acids and proteins are useful for diagnosing and treating carcinomas, for example lymphoma, breast cancer, prostate cancer or leukaemia, or for screening drug candidates or bloactive agents capable of binding to, or modulating the activity of, a carcinoma associated protein. The present sequence is the genomic DNA sequence of the human SYK gene which is a carcinoma associated gene of the invention.
                                                                                                                       Sequence
                                                                                                                                                                                                                                                                                                                                                                   New carcinoma associated nucleic acids and proteins, useful drug candidates, or for diagnosing and treating carcinomas, lymphoma, breast cancer, prostate cancer or leukemia.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              cancer diagnosis; cancer treatment; carcinoma; cytostatic; gene therapy;
lymphoma; breast cancer; prostate cancer; leukaemia; ds; human; SYK.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Human SYK gene genomic DNA sequence
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                                                                                                                                                                                                                                                                                                                                        Claim 1; SEQ ID NO 232; 793pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                   WPI; 2003-441462/41.
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                                                                                                                                                                                                                                                                                                         This invention relates to novel recombinant nucleic acids for use
                                                                                                                                                                                                                                                                                                                                                                                                                                                               Morris DW, Engelhard EK;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     WO2003039484-A2
                                                                          Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                cell. This sequence represents human cancer-associated genomic DNA of
57089
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 42048 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 42093
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                              CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 3116
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                                                                                                                       96594 BP; 27524 A; 20558 C; 21159 G;
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CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 57134
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                                                          Conservative
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Pred. No.
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Pred. No. 9.3e-11;
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9.4e-11;
                                                                                                                       26914 T;
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BXS
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                                                                      ADB72464
                                                                                          RESULT 102
                                                                                                                                                                                                                                                                                                                                                                                                                 The invention relates to recombinant carcinoma associated (CA) nucleic CC acid sequences from mouse and human (AbA01492-AbA03094), and to CC acid sequences from mouse expression vectors and host cells comprising a CC CA nucleic acid, a polypeptide (especially annibody) that specifically binds to the protein, and a biochip comprising CA nucleic acid or CC fragments thereof: The sequences of the invention were identified using CC oncogenic retroviruses, which insert into the genome of the host organism CC at random. Many of these do not carry transduced host oncogenes or CC pathogenic trans-acting viral genes, meaning that cancer incidence is a direct consequence of the effects of proviral integration into host CC protooncogenes. The CA nucleic acid sequences can be used to diagnose CC carcinoma (especially breast cancer, prostate cancer, lymphoma or CC leukaemia) or a propensity to carcinoma by determination of the sequence of a CA gene, or by determination of CA gene expression in particular CC tissues. CA nucleic acids, proteins and antibodies are also useful as CC present sequence represents a specifically claimed human CA nucleic acids sequence of the invention. Note: The complete sequence data for this complete sequence.
                                                                                                                                                                                                                                                            Query Match
Best Local S
Matches 46
ADB72464;
                                             ADB72464 standard; DNA; 96595 BP.
                                                                                                                                                                                                                                                                                                                                                          Sequence
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    New recombinant nucleic acid encoding carcinoma associated useful for preparing compositions for treating carcinomas.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   WPI; 2003-587068/55
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                gene;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Human; carcinoma associated; oncogene; carcinoma; cancer; breast;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human SYK carcinoma
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                                                                                                                                                                                                                                                                                      Local Similarity
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                                                                                                                                                                                                            3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 3116
                                                                                                                                                                 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 57135
                                                                                                                                                                                                                                                                                                                                                            96595 BP;
                                                                                                                                                                                                                                                            Conservative
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                                                                                                                                                                                                                                                            Mismatches
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                                                                                                                                                                                                                                                                                   DB 9; Le
                                                                                                                                                                                                                                                                                                          Length 96595;
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ID AD056
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Best Local Similarity
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08-NOV-2001;
30-NOV-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 The invention relates to a novel recombinant nucleic acid comprising a nucleotide sequence selected from any of the 660 sequences fully defined in the specification. A polynucleotide of the invention has cytostatic activity, and may have a use in gene therapy, or in a vaccine. The recombinant nucleic acids and polypeptides are useful for treating carcinoma, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and sarcomas. The present sequence represents a human gene of the invention.
                                                                                                                                                                    gene therapy; human; ds; gene; melanoma; melanoma associated polymorphic variation; SNP; single nucleotide polymorphism; cyclin-dependent
variation
                                                                                                                               Homo sapiens
                                                                                                                                                                                                                                                              Human cyclin-dependent kinase 10,
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ilarity 100.0%;
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"Single nucleotide polymorphism"
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variation 47754	/note= "Single nucleotide polymorphism"	variation
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variation 44692 *tag= au /note = scingle nucleotide	"Single nucleotide polymorphism"	variation
variation /*tag= at /note= "Single nucleotide		variation
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variation	/*tag= t /note= "Single nucleotide polymorphism"	;
variation 43164	/"Cag= s /note= "Single nucleotide polymorphism" 18077	variation
variation 42477 /*tag= aq	/note= "Single nucleotide polymorphism" 17564	variation
	16824 /*tag= r	Variation
variation 41955	/*tag= q /note= "Single nucleotide polymorphism"	
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variation 40292 /*tag= al	/note= "Single nucleotide polymorphism" 14845	variation
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variation 39762	/note= "Single nucleotide polymorphism"	
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variation 34152	/"tag=) /note= "Single nucleotide polymorphism"	
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variation 27859	/note= "Single nucleotide polymorphism" 7960	variation
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23 standard; DNA; 99250 BP. 23; 23; -2005 (first entry) cyclin-dependent kinase 10 (CDX10) genomic DNA. ma; DNA polymorphism; SNP detection; cytostatic; gene therapy; SNP; nucleotide polymorphism; gene; ds; chromosome 16. Location/Qualifiers /*tag= a /*tag= b /*tag= b /*tag= b /*tag= c /*tag= c /*tag= c /*tag= c /*tag= d /*standard_name= "Single nucleotide polymorphism" End End End End End End End En	1.5%; Score 46; DB 12; Length 99100; imilarity 100.0%; Pred. No. 9.3e-11; Conservative 0; Mismatches 0; Indels 0; Gaps 0; CAAGATTGTGCCACTCCAGCCTGGGCAACAGAGCAAGACTC 3116 CAAGATTGTGCCACTCCAGCCTGGGCAACAGAGCAAGACTC 16476 EAAGATTGTGCCACTCCAGCCTGGGCAACAGAGCAAGACTC 16476	/*tag= ay //note= "Single nucleotide polymorphism" /*tag= az // 49672 /*tag= az // 50476 /*tag= ba // 50476 /*tag= ba // 50525 /*tag= bb // 50621 /*tag= bb // 50621 /*tag= bc // 50621 /*tag= bd // 50622 /*tag= bf // 50622
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Best Local Similarity
Matches 46; Conserv
                                                                            New isolated cancer associated nucleic contiguous nucleotides, useful for dia cancers such as leukemia and lymphoma.
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present invention relates to cancer associated sequences (ADQ97025-

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RESULT 107
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Sequence split into 5
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 109661 BP; 30680 A; 19371 C; 19986 G; 35350 T; 0 U; 4274 Other;
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                                        ADZ13631;
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ADG70447 3
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9.3e-11;
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Diagnosis; DNA microarray; microarray; biochip; cancer; neoplasm; cytostatic; gene; ds.
Morris DW,
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                                                                                                                                                                                          WO2005031001-A2
                                      (CHIR ) CHIRON CORP
Malandro
                                                                                                                2004WO-US031617
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Nucleic acid array useful for detecting cancer associated nucleic acid, comprises two or more nucleic acid probes.

WPI; 2005-273395/28

Disclosure; SEQ ID NO 1151; 198pp; English.

CC The invention also relates to a peptide array comprising two or more isolated polypeptides encoded by a CA nucleic acid sequence, a compound CC that binds to a polypeptide, an isolated antibody or its fragment which CC binds to a polypeptide, which is prepared by immunizing a host animal CC with a composition comprising the polypeptide or its antigen binding CC fragment and collecting cells from the host expressing antibodies against the antigen or its antigen binding fragment, a composition comprising the CC antibody and a carrier, a method of screening for anticancer activity, a CC method of detecting a CA nucleic acid, a method of diagnosing cancer, a CC method of tracting cancer and a method of inhibiting expression of a CA concleic acid in a cell. The CA nucleic acids are useful for detecting CA cC nucleic acids. The antibody is useful for detecting the presence or cabsence of cancer cells in an individual which involves contacting cells from the individual with the antibody and detecting a complex of a CA composed from the cancer cells in an individual which involves contacting cells in an individual. The composition is useful for inhibiting growth of cancer cells in an individual. The presence of cancer cells in the cancer cells in an individual or for delivering a therapeutic agent to cancer cells in an individual. The invention is also useful for diagnosing concer, for treating cancer and for inhibiting expression of a CA gene in a cell. This sequence represents human cancer-associated genomic DNA of associated (CA) nucleic acid, comprising two or more nucleic relates to a nucleic acid array for detecting a cancer probes

Sequence 420555 BP; 131028A; 77271C; 78657G; 131031T; 0U; 25680ther;

Query Match Best Local S Matches 46

Local Similarity nes 46; Conserv

larity 100.0%; Conservative (

1.5%; Score 46; DB 100.0%; Pred. No. 9: 1ve 0; Mismatches

DB 14; 9.3e-11; 0;

Length 110000; Indels

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AXUABABBBB RESULT 109
ADZ13620_0
WP Sequence split into 밁 S ADZ13620 standard; Fragment Name ADZ13620 0 ADZ13620 1 ADZ13620 2 ADZ13620 3 ADZ13620 3 50592 3077 TGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 50637 TGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122 ທ DNA; 420555 fragments 100001 200001 300001 400001 Begin 110000 210000 310000 410000 420555 BP. End LOCUS ADZ13620 Accession Adz13620

Human cancer-associated genomic DNA #99

ADZ13620;

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CC The invention also relates to a peptide array comprising two or more consisted polypeptides encoded by a CA nucleic acid sequence, a compound content and the polypeptide, which is prepared by immunizing a host animal content and collecting cells from the host expressing antibodies against the antigen or its antigen binding fragment, a composition comprising the content and collecting cells from the host expressing antibodies against the antipen or its antigen binding fragment, a composition comprising the content and collecting cells from the host expressing antibodies against the content of the composition comprising the content of the composition comprising the content of the carting and a method of diagnosing cancer, a composition of a CA nucleic acid in a cell. The CA nucleic acids are useful for detecting CA concelic acids in a cells in an individual which involves contacting cells content from the cancer cells in an individual which involves contacting cells content from the cancer cells and the antibody, where the detection of the complex correlates with the presence of cancer cells in the composition is useful for inhibiting growth of cancer cells in an individual. The invention is also useful for dagnosing concer, for treating cancer and for inhibiting expression of a CA gene in a cell. This sequence represents human cancer-associated genomic DNA of the invention is also useful for dagnosing.
P Sequence split into 4

P Fragment Name
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                                                                                                                                                         Query Match
                                                                                                                                                                                                                                                                                 The invention relates to a method of Identifying p53 pathway modulating agent. The method involves contacting a test agent with an assay system comprising a purified cation C1- cotransporter interactor protein (CIP) polypeptide or polynucleotide, or their functionally active fragment or derivative. The method is useful for identifying modulators of the p53 pathway particularly for identifying agents for treating disorders (e.g. breast cancer) associated with defective p53 function. Modulators of the invention are useful as targets for novel therapeutics. CIP sequences are useful as modifiers of the p53 pathway, and as therapeutic targets for apoptotic or cell proliferation disorders. The invention is useful in comparator accounts.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             05-JUN-2001; 2001US-0296076P.
10-OCT-2001; 2001US-0328605P.
15-FEB-2002; 2002US-0357253P.
                                                                                                                                                                                                                 Sequence 117962 BP; 27840 A; 32096 C; 30624 G;
                                                                                                                                                                                                                                                              gene therapy. The present sequence is human CIP DNA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Identifying p53 pathway modulators for treating or diagnosing d with defective p53 function e.g. breast cancer, by providing an system having a purified cotransporter interactor protein (CIP) polypeptide or nucleic acid.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       03-JUN-2002; 2002WO-US017473.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; p53 pathway; chloride cotransporter interactor protein; CIP; angiogenic disorder; cell proliferation disorder; apoptotic disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Human CIP DNA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       (EXEL-) EXELIXIS INC.
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                                                                                                                                госат
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                                                     3071
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  cancer; gene therapy; ds.
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CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 18443
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                                                  CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC
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                                                                                                       Conservative 0;
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                                                                                                                                                            1.5%;
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                                                                                                    Score 46; DB; Pred. No. 9.3
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              365720
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                                                                                                                                DB 8; Le
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DB 14; I
9.3e-11;
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                                                                                                                                                                                                                 27402 T; 0 U; 0
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RESULT 113
ADC8762/C
ID ADC876
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ACN43862
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The present invention relates to novel DNA and protein sequences which care associated with carcinomas. The sequences are useful for: (1) for screening drug candidates; (11) for screening of bioactive agent capable of modulating the activity of CAP; (11) for screening of a bioactive agent capable of modulating the activity of CAP; (11) for carcinoma; (vi) for inhibiting the activity of CAP; (11) for treating carcinoma; (vii) for neutralizing the effect of CAP; (vii) for treating carcinoma; (viii) for neutralizing the effect of CAP; (vii) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma associated (CA) gene copy number. In addition, the CAP genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent CAP which no sequence data was published
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Query Match
Best Local
                                                           ds; human; GPCR; guanosine triphosphate-binding protein coupled receptor;
                                                                                                                       Human GPCR related polynucleotide
                                                                                                                                                                                  01-JAN-2004
                                                                                                                                                                                                                                                                                                      ADC87620 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 141463 BP; 40336 A; 28306 C; 29237 G; 43584 T; 0 U; 0
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Recombinant nucleic acid useful for diagnosis comprises a nucleotide sequence.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                       73703
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                               therapy.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Similarity
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ilarity 100.0%;
Conservative (
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                                                                                                                                                                                                                                                                                                      DNA; 144792
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Pred. No. 9.2e-11
0; Mismatches 0
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                                                                                                                       SEQ ID NO:2073.
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RESULT 114
ADL13904/c
ID ADL13904 standard; DNA; 164772
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                                                                                                                                                                                                                                    ds; gene; osteopathic; antiinflammatory; antiarthritic; gene therapy; joint space narrowing; osteophyte development; joint pain; osteoarthritis; SNP; single nucleotide polymorphism.
             WPI; 2003-559141/52
                                      Jones KA,
                                                                                                                                                                                                           Homo sapiens.
                                                                                                                                                                                                                                                                                            Osteoarthritis-associated polymorphic nucleotide #436.
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                                                                                              20-DEC-2001; 2001US-0342603P
                                                                                                                           19-DEC-2002; 2002WO-US041225
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(ADSC-)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                         60793
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CENT ADVANCED SCI & TECHNOLOGY INCUBATI
                                                                    INCYTE GENOMICS
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                                        Schafer A;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   the invention.
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                                                                                                                                                                                                                                                                                                                        (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Aburatani H;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Mismatches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             9.2e-11;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                         60748
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of the
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Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding Disclosure; SEQ ID NO 436; 297pp; English

individual to joint space narrowing and/or osteophyte development and/or joint pain comprising identifying whether the individual has at least one polymorphism in a polymuclectide encoding at least one of the protein listed in the specification. The methods, composition and agent are useful for modulating the susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain that is associated with a disease, preferably osteoarthritis. The cell line and the non-human animal are useful for screening for an agent for diagnosing an individual having susceptibility to joint space narrowing and/or osteophyte development and/or joint space narrowing and/or osteophyte development and/or joint pain. This sequence corresponds to the polymucleotide encoding a protein listed in the specification. (Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences). invention relates to a method of determining susceptibility of an

Sequence 164772 BP; 50645 A; 32137 C; 31960 G; 50022 T; 0 U; 8 Other;

Matches Query Match Best Local Similarity Conservative 1.5%; 0 Score 46; Pred. No. Mismatches DB 10; 9.2e-11; .. Length 164772; Indels 0 Gaps 0

용 δ 111876 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 111831 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940

RESULT 115 ACN44262

ACN44262 standard; DNA; 168821 ВP

ACN44262

18-NOV-2004 (first entry

Human genomic sequence hCG18035.

Cytostatic; carcinoma; lymphoma; cancer; human; gene; 88

Homo

WO2003073826-A2

12-SEP-2003

28-FEB-2003; 2003WO-US006235.

01-MAR-2002; 2002US-00087192.

(SAGR-) SAGRES DISCOVERY

Morris

2003-328604/31.

Recombinant nucleic acid useful for diagnosis and comprises a nucleotide sequence. treatment о f carcinoma

Claim 1; SEQ

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NO 622; Opp; English.

The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bloactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bloactive agent capable of modulating the activity of CAP; (Iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing

RESULT 116
ADL13935
AC ADL139
XX ADL139
XX DT 06-MAY
XX Ods; ge
KW dds; ge
KW joint
XX Joint
XX Osteoa
XX SSSSSSSXX 밁 S Best Query Match carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating carcinoma; (vii) for neutralizing the effect of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent US2002182586AI, for which no sequence data was published Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polymucleotide encoding ds; gene; osteopathic; antiinflammatory; antiarthritic; joint space narrowing; osteophyte development; joint pai Sequence 168821 BP; Jones KA, 03-JUL-2003 Homo sapiens osteoarthritis; Osteoarthritis-associated polymorphic 06-MAY-2004 ADL13935; ADL13935 standard; DNA; 177866 WPI; 2003-559141/52 20-DEC-2001; 2001US-0342603P 19-DEC-2002; 2002WO-US041225 WO2003054166-A2 protein. 161763 Local 2895 46; Similarity INCYTE GENOMICS GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 161808 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT Schafer A; Conservative (first entry) SNP; 39588 A; 43389 C; 45655 G; 40189 T; 0 U; 0 single 100.0%; 1.5%; nucleotide polymorphism. 0 Score 46; Pred. No. BP Mismatches nucleotide DB 11; 9.1e-11 .1e-11; joint pain; Length 168821; Indels gene 2940 0, Other; Gaps 0

Disclosure; SEQ ID NO 467; 297pp; English.

The invention relates to a method of determining susceptibility of an CC individual to joint space narrowing and/or osteophyte development and/or CC joint pain comprising identifying whether the individual has at least one CC polymorphism in a polymuclectide encoding at least one of the protein CC listed in the specification. The methods, composition and agent are CC useful for modulating the susceptibility of an individual to joint space CC narrowing and/or osteophyte development and/or joint pain that is CC associated with a disease, preferably osteoarthritis. The cell line and CC in individual having susceptibility to joint space narrowing and/or control to control to the polymuclectide encoding a protein listed in the specification. (Note: CC che polymuclectide encoding a protein listed in the specification but was obtained in electronic format directly from WIPO at CC ftp.wipo.int/pub/published_pct_sequences).

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Sequence 177866 BP;

53227 A;

36632 C; 36825 G; 51154 T; 0 U; 28

Other;

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RESULT 118
ABQ75562
ID ABQ755
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AC ABQ755
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AC ABQ755
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Best Local
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                                                                                                                                                                                                                                                                               The present invention describes the human SLC5AB protein (I), which is a cell surface protein. (I) has cyrostatic activity, and can be used in gene therapy. (I) can be used in detecting and treating SLC5AB sassociated cancer, e.g. colon cancer, breast cancer, thyroid cancer or stomach cancer. (I) is also useful in screening assays, predictive medicine and in diagnostic and prognostic assays. The human SLC5AB gene is located on chromosome 12. The present sequence is used in the exemplification of the
  11-NOV-2002
                             ABQ75562;
                                                                                                                                                                                                                                         Sequence 181257 BP; 53237 A; 35656 C; 35971 G; 56393 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                               Claim 6; SEQ ID NO 2; 207pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                         New SLC5A8 polypeptide, useful for detecting and treating SLC5A8-associated cancer, e.g. colon, breast, thyroid or stomach cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Markowitz
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      05-JUN-2002; 2002US-0386653P
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            human; SLC5A8; cell surface protein; cytostatic; gene therapy; SLC5A8-associated cancer; colon cancer; breast cancer; thyroid cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Human SLC5A8 gene SEQ ID NO:2.
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                                                       ABQ75562 standard;
                                                                                                                                                                                                                                                                     present invention.
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                                                                                                                                                                                                Local Similarity
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                                                                                                                            GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 99645
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                                                                                                                                                                                   Conservative
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                                                       DNA;
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                                                       188888
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Pred. No.
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9.1e-11;
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Cloning; characterisation; human; cytochrome P450; CYP 27C1; cytostatic; thyromimetic; antidiabetic; antipsoriatic; tuberculostatic; osteopathic; dermatological; antilipaemic; gene therapy; vaccine; Vitamin D; diabetes; vitamin D metabolite deficiency; hyperparathyroidism; hypoparathyroidism; medullary carcinoma; psoriasis; sarcoidosis; tuberculosis; osteomalacia; chronic renal disease; vitamin D dependent rickets; anticonvulsant; fibrogenesis imperfecta ossium; osteotitis fibrosa cystica; osteoporosis; osteopaenia; osteosclerosis; renal osteodystrophy; rickets; steatorriboea; glucocorticoid antagonism; idiopathic hypercalcaemia; tropical sprue; malabsorption syndrome; cholesterol steroid; lipid metabolic disorder; cene; de:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human related CYP 27C1 clone RP11-30F3 SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               NO:21.
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09-FEB-2001; 2001US-0267410P 22-AUG-2002. WO200264765-A2 11-FEB-2002; 2002WO-CA000163.

Wisniewski

WPI; 2002-657595/70

New nucleic acid molecules encoding cytochrome P450 proteins, human CYP 27C1 and a hybrid homologs from Xenopus laevis, useful for treating diseases related to vitamin D or vitamin D metabolite deficiency, e.g. parathyroidism and diabetes.

Example 1; Fig 1A; 209pp; English.

The present invention describes an isolated nucleic acid molecule (I) CC encoding human cytochrome P450, CPP 27C1, and a hybrid homologue from CC Xenopus laevis. (I) has thyromimetic, antidiabetic, cytostatic, cc antilpsoriatic, tuberculostatic, osteopathic, dermatological and cantilpsoriatic acid molecules, and can be used in gene therapy and in vaccines. CC antilpsoriation are useful for treating diseases related to vitamin D or vitamin D metabolite deficiency, e.g. hyper- and hypo-parathyroidism, CC osudohypo-parathyroidism, Secondary hyperparathyroidsm, diabetes, cc disease, hypophosphatemic VDRR, vitamin D dependent rickets, anticonvulsant treatment, fibrogenesis imperfecta ossium, osteoititis cfibrosa cystica, osteomalacia, osteoporosis, osteopania, osteosclerosis, renal osteodystrophy, rickets, glucocorticoid antagonism, idiopathic cor cholesterol, steroid and other lipid metabolic disorders. The present cc which is given in an example from the present invention

Sequence 188888 BP; 51055 A; 42661 C; 43560 G; 47708 T; 0 U; 3904 Other;

0

RESULT 119 ADL13570/c ID ADL135 XX AC ADL135 밁 Ş Query Match
Best Local Similarity
Matches 46; Conserva ADL13570 standard; DNA; 193672 BP 142425 2895 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACAT 2940 Conservative 100.0%; 1.5%; Score 46; 100.0%; Pred. No. Mismatches DB 6; 9.1e-11; Length 188888; Gaps

ADL13570;

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Best Local S
Matches 46
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ds; gene; osteopathic; antiinflammatory; antiarthritic; gene therapy; joint space narrowing; osteophyte development; joint pain; osteoarthritis; SNP; single nucleotide polymorphism.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               20-DEC-2001; 2001US-0342603P
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               Homo sapiens
                                               Cytostatic; carcinoma; lymphoma; cancer; human; gene;
                                                                                Human genomic
                                                                                                                 18-NOV-2004
                                                                                                                                                    ACN44650
                                                                                                                                                                                  ACN44650 standard; DNA; 256157 BP
                                                                                                                                                                                                                                                                                                                                                                                                      Sequence 193672 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                      ftp.wipo.int/pub/published_pct_sequences).
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                                                                                                                                                                                                                                                                                                                                                    Local Similarity
                                                                                                                                                                                                                                                                     3015 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 2970
                                                                                                                                                                                                                                                                                                   3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTC 3116
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             SEQ ID NO 102;
                                                                                                                                                                                                                                                                                                                                    Conservative 0;
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                                                                                                                 (first entry)
                                                                                sequence hCG38672.
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                                                                                                                                                                                                                                                                                                                                                                       1.5%;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          297pp; English.
                                                                                                                                                                                                                                                                                                                                                      Score 46; Pred. No.
                                                                                                                                                                                                                                                                                                                                 Mismatches
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9.1e-11;
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RESULT 121
ABD3357
XX ABD335
XX ABD335
XX ABD335
XX INIMAL
DT 18-NOV
XX Human;
XW Human;
XW Homo 6
XX Homo 6
XX Homo 6
XX INIMAL
PN WO2004
XX 15-DE(
XX 17-DE(
XX INIMAL
PR 17-DE(
XX INIMAL
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Matches 46; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 256157 BP; 70370 A; 54568 C; 55511 G; 73304 T; 0 U; 2404 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.
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                                                                     Morris DW,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human cancer-associated (CA) gene HD07-114.
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                                                                                                                                                                                               17-DEC-2002; 2002US-00322281
                                                                                                                                                                                                                                                           15-DEC-2003; 2003WO-US040081.
                                                                                                                                                                                                                                                                                                                                                                                            WO2004058146-A2
                                                                                                                                                                                                                                                                                                                                                                                                                                                    Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Human; cancer-associated protein; CAP; cancer-associated gene; CA; gene;
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                                                                                                                                    (SAGR-) SAGRES DISCOVERY INC
                                                                                                                                                                                                                                                                                                                             15-JUL-2004.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      2895
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                                                                     Malandro MS;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              1.5%; So lilarity 100.0%; I conservative 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   cytostatic.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Score 46;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Mismatches
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9e-11;
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WPI; 2004-499109/47

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    associated (CA) nucleic acids encoding them. The invention also relates to a method for treating cancers involving administering to a patient an inhibitor of CAP, and a method of screening for anticancer activity in a potential drug involving providing a cell that expresses a CA gene, contacting a tissue sample derived from a cancer cell with an anticancer drug candidate and monitoring the effect of the anticancer drug candidate on expression of the CA gene. The CAP proteins are useful for detecting cancer associated with expression of a CAP protein in a test cell sample and for screening for a bioactive agent capable of modulating the activity of a CAP protein. The CAP nucleic acids are useful for diagnosing cancer, involving determining the expression of a CA nucleic acid in a tissue. This sequence represents a human CA gene of the invention. Note:

The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO
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Best Local S
Matches 46
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are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bioactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bioactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
                                                                                                                      Claim
                                                                                                                                                                      Recombinant nucleic acid useful for diagnosis and treatment
                                                                                                                                                                                                                                                                                                         01-MAR-2002; 2002US-00087192
                                                                                                                                                                                                                                                                                                                                          28-FEB-2003; 2003WO-US006235.
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                                                                                                                                                                                                                                                                                                                                                                                                                                               Homo
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Cytostatic; carcinoma;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human genomic sequence hCG17121.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    18-NOV-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ACN44350;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Sequence 256157 BP; 70370 A; 54568 C;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Novel human cancer associated protein encoded within open reading frame of cancer associated gene, useful as targets for diagnosing cancer.
                                                                                                                                                                                                                                                                         (SAGR-) SAGRES DISCOVERY.
                                                                                                                                                                                                                                                                                                                                                                            12-SEP-2003.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   invention relates to cancer-associated proteins (CAP) and the cancer
                                                                                    present invention relates to novel DNA and protein sequences which
                                                                                                                                                                                                       2003-328604/31
                                                                                                                                                                                                                                                                                                                                                                                                                                               sapiens.
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                                                                                                                                                    a nucleotide sequence.
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                                                                                                                      ID NO 754;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  (first
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                               lymphoma; cancer; human; gene;
                                                                                                                   Opp; English.
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Pred. No.
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9e-11;
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The present invention describes human cancer-associated (CA) nucleotide compenses (I). Also described: (1) an expression vector comprising (I); (C) a host cell comprising (I) or the expression vector; (3) a microarray comprising a CA nucleic acid; (4) an isolated polypeptide encoded comprising a CA nucleic acid; (4) an isolated polypeptide encoded comprision frame of a CA sequence; (5) an isolated antibody, (6) or its antigen binding frame of a CA sequence; (5) an isolated antibody, (6) comprision frame of a CA sequence; (5) an isolated antibody, (6) comprision frame of a CA sequence; (5) an isolated above; (7) a planmaceutical composition comprising the antibody and a pharmaceutical composition comprising cancer cells, comprising che above (monoclonal) antibody or polynucleotide that selectively comprising the above (monoclonal) antibody or polynucleotide that selectively (9) comprisions for diagnosing cancer or for detecting the presence or absence of concer cells in an individual; (11) a method for inhibiting growth of concer cells in an individual; (11) a method for delivering a therapeutic agent to cancer cells in an individual; (11) an electronic library comprising the polynucleotide or polypeptide, or their fragments,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           RESULT 123
ADQ59440/c
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Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent US2002182586A1, for which no sequence data was published
                                                                                                                                                                                                                                                                                                                                                                                                New cancer-associated nucleic acid for diagnosing, preventing cancer (e.g. lymphoma) or for screening agents that may be use treating or preventing cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        human; cancer-associated gene; cancer-associated protein; cytostatic; gene therapy; vaccine; tyrosine kinase antagonist; G-protein coupled receptor antagonist; cancer; lymphoma; gene; ds.
                                                                                                                                                                                                                                                                                                                                                              Claim 16;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                ADQ59440;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             212141 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGG 212186
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                                                                                                                                                                                                                                                                                                                                                              SEQ ID NO 76;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Malandro
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                                                                                                                                                                                                                                                                                                                                                         143pp; English.
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Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            gene sequence SEQ ID NO:76
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Query Match
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  in the exemplification of the present inverted for this patent did not form part of the pobtained in electronic format directly from the property from the property of the present directly from the property of the present directly from the property of the present directly from the present directly di
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AAS27638 standard; DNA; 145
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; 2000US-0179065P.

2000US-0184664P;

2000US-0184659P;

2000US-0186350P;

2000US-0189874P;

2000US-0199123P;

2000US-0298159;

2000US-0205515P;

2000US-0219467P;

2000US-0215135P;

2000US-0216647P;

2000US-0217487P;

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Conservative 0;
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            The invention relates to novel isolated polypeptides (I), and polynucleotides (II). (II) and the antibody to (I) are useful for diagnosing, preventing and treating diseases including immune system disorders (e.g. congenital and acquired immunodeficiencies, autoimmune disorders (e.g. rheumatoid arthritis), inflammatory conditions, organ transplant rejections and graft versus host disease, infectious diseases (e.g. hepatitis C), bleeding disorders, haemoglobin abnormalities and other blood-related disorders (sickle cell anaemia), myeloproliferative disorders, primary haematopoietic disorders, hyperproliferative disorders (e.g. Gaucher's disease, Parkinson's disease), chromosomal abnormalities (Down syndrome), ischaemic injury (e.g. stroke), renal disorders (e.g.
                                                                                                                                                                                                                                                                                                           Novel polypeptides useful for diagnosing, treating, prevprognosing disorders related to the proteins, including disorders and neuronal disorders.
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(Down syndrome), ischaemic injury (e.g. stroke), renal disorders glomerulonephritis), cardiovascular disorders (e.g. arrhythmia),
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respiratory disorders, dermatological disorders, in wound healing, epithelial cell proliferation, endocrine disorders (e.g. Addison's disease), reproductive system disorders, gastrointestinal disorder (inflammatory disorders), liver disorders (cirrhosis), as stimulators of B-cell responsiveness to pathogens, activators of T-cells, to induce higher affinity antibodies, and as a means to induce tumour proliferation in pathologies e.g. acquired immune deficiency syndrome (ADIS), AA26976-AAS27850 represent novel signal transduction pathway protein coding sequences and PCR primers of the invention
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Sequence 145 BP; 13 A; 36 C; 21 G 75 T; 0 U; 0 Other;

Query Match Local 3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122 120 ch 1.4%; Solution 1.4 Score 45; DB Pred. No. 3.4 0; Mismatches 3.4e-10; hes 0; DB 4; Length Indels 145; 76 0 Gaps 0

06-NOV-2001 AAK68506; AAK68506 standard; DNA; 145 (first entry) ₽P

Human immune/haematopoietic antigen genomic sequence SEQ ID NO:23318

cytostatic; Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; gene therapy; vaccine; metastasis; ds.

WO200157182-A2 sapiens.

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31-JAN-2000; 04-FEB-2000; 24-FEB-2000; 02-MAR-2000; 16-MAR-2000; 17-MAR-2000; 18-APR-2000; 19-MAY-2000; 07-JUN-2000; 2000US-0186628P.
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                                                                                                                                                                           amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic cactivity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by recifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the curleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, cancers and treat immune/haematopoietic-derived cells. AAK64703 cc cancers and cancer metastases of haematopoietic antigen genomic to AAK647694 represent human immune/haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention
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08-DEC-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         antito acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, cancers and cancer metastases of haematopoietic-derived cells. AAK64703
          Central nervous system; CNS; autoimmune disease; rheumatoid arthritis; hyperproliferative disorder; neoplasm; cardiovascular disorder; cardiac arrest; cerebrovascular disorder; ischaemia; angiogenesis; nervous system disorder; Alzheimer's disease; AIDS; ocular disorder; acquired immunodeficiency virus; dysphagia; gastrointestinal disorder; adenocarcinoma; reproductive system disorder; testicular feminisation; endocrine disorder; diabetes; cancer; leukaemia; neovascularisation; respiratory disorder; renal disorder; testicular feminisation; respiratory disorder; wound healing; cell proliferation; skin aging; food additive; food preservative; gene therapy; gene; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          08-DEC-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Disclosure;
                                                                                                                                                                                                                                                 Genomic
                                                                                                                                                                                                                                                                                                                                                                        ABK44042 standard; DNA; 145 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 145
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AAK87694 represent human immune/haematopoietic antigen genomic quences from the present invention. AAK54942 to AAK54950 and AAM82169 present sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   26
                                                                                                                                                                                                                                               DNA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                               GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            to AAK64702 encode the human immune/haemat id sequences given in AAM82170 to AAM91921.
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2000US-0251719P.
2000US-0251479P.
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2000US-0251989P.
2000US-0254097P.
2001US-0259678P.
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                                                                                                                                                                                                                                                 encoding novel central nervous system protein #244.
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100.0%; Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     3.4e-10;
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07-JUN-2000;
28-JUN-2000;
30-JUN-2000;
30-JUN-2000;
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18-APR-2000;
19-MAY-2000;
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2000US-0199076P

2000US-0190076P

2000US-0205515P

2000US-02151886P

2000US-0216647P

2000US-0217487P

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             2000US-0232401P.
2000US-0233063P.
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2000US-0234273P.
2000US-0234273P.
2000US-0234279P.
2000US-0234299P.
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2000US-0229513P.
2000US-0230437P.
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2000US-0231242P.
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2000US-0184664P.
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2000US-0232400P.
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2000US-0231414P.
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27-SEP-2000; 27-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000;

preventing, additives o

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2000US-0250160P.
2000US-0250391P.
2000US-02510391P.
2000US-0251988P.
2000US-0251479P.
2000US-0251866P.
2000US-0251866P.
2000US-0251869P.
2000US-0251869P.
2000US-0251990P.
2000US-0254997P.
2001US-0259678P.
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2000US-0237038P
2000US-0237039P
2000US-0237040P
2000US-0239935P
2000US-024122IP
2000US-0241785P
2000US-0241786P
2000US-0241808P
2000US-0241808P
2000US-024671P
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2000US-0235836P.
2000US-023632PP.
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     GENOME SCI
ARESULT 128
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Best Local S
Matches 45
  31-JAN-2000;
04-FEB-2000;
28-JUN-2000;
07-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                      ds; gene; human; autoimmune disease; Parkinson's disease; silicosis; gastrointestinal disease; atherosclerosis; haemophilia; thrombocytopenia; immunosuppressive agent; adjuvant; enhance immune response; higher affinity antibody induction; increased serum immunoglobulin concentration.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    angiogenesis, nervous system disorders e.g. Alzheimer's disease and amylotrophic lateral sclerosis, infections caused by bacteria, viruses e.g. Acquired immunodeficiency virus (AIDS) and fungi, ocular disorders e.g. dysphagia, e.g. corneal infection, gastrointestinal disorders e.g. dysphagia, adenocarcinomas and irritable bowel syndrome, reproductive system disorders e.g. testicular feminisation, endocrine disorders e.g. diabete and pituitary dwarfism, cancers and disorders at the cellular level e.g.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            leukaemia, disorders involving neovascularisation e.g. malignancies, respiratory disorders e.g. nonallergic rhinitis, renal disorders e.g. acute kidney failure and blood related disorders e.g. myocardial infarction. The polypeptides can also be used to aid wound healing and epithelial cell proliferation, to prevent skin aging due to sunburn, to maintain organs before transplantation, for supporting cell culture of primary tissues, to regenerate tissues and in chemotaxis. The
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       04-DEC-2003
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ADB94441
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 carbohydrate,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  pathological condition. Disorders which are diagnosed or treated include autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative disorders e.g. neoplasms of the breast or liver, cardiovascular disorders
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Disclosure; SEQ ID NO 1230;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    New isolated nucleic acid encoding a protein for diagnosing, treating or ameliorating medical conditions and used as food
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Rogen
                                                                                                                                          17-JAN-2001;
                                                                                                                                                                                                14-NOV-2002.
                                                                                                                                                                                                                                                 US2002168711-A1
                                                                                                                                                                                                                                                                                                        Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Novel human protein DNA #50.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              ADB94441;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        polypeptides can also be used as a food additive or preservative to increase or decrease storage capabilities, fat content, lipid, proto
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     isorders e.g. neoplasms of the breast or liver, cardiovascular disorde.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischaemia,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  3078
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         45;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
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Pred. No.
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17-NOV-2000; 17-NOV-2000; 17-NOV-2000; 17-NOV-2000;

17-NOV-2000; 17-NOV-2000;

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useful for diagnosing a pathological condition or a susceptibility to a pathological condition in a subject, by determining the presence or amount of expression of the polypeptide in a biological sample and diagnosing a pathological condition or a susceptibility to a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide. The polypeptide is also useful for identifying a binding partner to the polypeptide, which involves contacting the polypeptide with a binding partner and determining whether the binding partner effects an activity of the polypeptide. The polypeptide or the mucleic acid encoding the polypeptide is useful for preventing, treating, or ameliorating a medical condition, which involves administration.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        11-JUL-2000

11-JUL-2000

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26-JUL-2000

26-JUL-2000

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10-SEP-2000

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                                                                                                                                                                                                                                                                                                                                     New isolated polypeptide useful for diagnosing and treating immunosuppressive conditions such as autoimmune disease and
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(RUBE/)
(BARA/)
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                                                                                                                                                                                                                                                                               Disclosure;
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RUBEN S
BARASH
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2000US-0225268P.
2000US-0225757P.
2000US-02354274P.
2000US-0234274P.
2000US-0234274P.
2000US-0235834P.
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2000US-0236367P.
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2000US-023703P
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Parkinson's

18-APR-2000

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AD1550

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                             31-JAN-2000;
04-FEB-2000;
04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            immune system disorder; diabetes; rheumatoid arthritis; systemic lupus erythematosus; autoimmune thyroiditis; haemolytic anaen inflammatory disorder; ischaemia-reperfusion injury; inflammatory bowel disease; Crohn's disease; infectious disease; HIV infection; hepatitis infection; bacterial infection; fungal infection; parasitic infection; muscular disorder; reproductive disorder; gastrointestinal disorder; pulmonary disorder; cardiovascular disorder; atherosclerosis; arrhythmia, myocarditis; renal disorder; acute glomerulonephritis; pyelonephritis; human; derenal lithiasis; proliferative disorder; cancerous diseases; human; de
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    anti-HIV; hepatotropic; virucide; anti-HIV; hepatotropic; virucide; anti-HIV; hepatotropic; virucide; anti-Acteria; fungicide; anti-Acteria; empiratory; antiparasitic; muscular; gynaecological; gastrointestinal; respiratory; cardiovascular; antiarteriosclerotic; antiarrhythmic; cardiant; nephrotropic; litholytic; cytostatic; gene therapy; neural disorder; Alzheimer; disease; Parkinson's disease; Huntington's chorea; amyotrophic lateral sclerosis; multiple sclerosis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                neuroprotective; nootropic; antiparkinsonian; anticonvulsant;
antidiabetic; antirheumatic; antiarthritic; dermatological;
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45; Conserv
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2000US-0180664P.
2000US-0184664P.
2000US-0186350P.
2000US-018974P.
2000US-0199076P.
2000US-0198123P.
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Pred. No. 3.4
0; Mismatches
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3.4e-10;
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20-OCT-2000

20-OCT-2000

20-OCT-2000

20-OCT-2000

01-NOV-2000

08-NOV-2000

17-NOV-2000

17-NO
                                                                       New polypeptides and nuc
preventing, diagnosing,
conditions e.g. neural c
diseases.
                                                                                                                                                                                                                                                (ROSE/)
(RUBE/)
(BARA/)
                                    Disclosure;
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RUBEN S
BARASH
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2000US-0241785P.
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2000US-0248471P.
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2000US-0246471P.
2000US-0246475P.
2000US-0246478P.
2000US-0246523P.
2000US-0246523P.
2000US-0246528P.
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2000US-0246610P.
2000US-0246611P.
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2000US-0249209P.
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2000US-025929P.
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                                                                                                                                                                                                            SM,
                                  NO 1230; 413pp; English
                                                                                         l nucleic acid molecules, useful for detecting,
.ng, prognosticating, treating or ameliorating medi.
ral disorders, reproductive disorders or infectious
                                                                                                                                                                                                                Barash
                                                                                                                                                                                                                SC
                                                                                                                medical
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The

invention describes an isolated polypeptide comprising

an

amino acid

19-MAY-2000
28-JUN-2000
28-JUN-2000
28-JUN-2000
29-JUN-2000
11-JUL-2000
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14-AUG-2000
11-SEP-2000
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2000US-020515P.
2000US-0214886P.
2000US-0214886P.
2000US-0214887P.
2000US-0216880P.
2000US-0216880P.
2000US-021513P.
2000US-02152964P.
2000US-0224519P.
2000US-0225214P.
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2000US-0225213P.
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XX Human;
XX Homo g
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XX US2003
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XX (DRMA)
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The invention relates to an isolated polymucleotide comprising any one o 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was determined by the technique of SBH (sequencing by hybridisation). Also included is a purified polypeptide comprising a sequence corresponding t a reading frame of the novel polymucleotide. The nucleic acid sequences are useful in diagnostics as expressed sequence tags (EST) for identifying expressed genes or for physical mapping of the human genome, in forensics, in assessing biodiversities, or in identifying mutations responsible for genetic disorders and other traits. The nucleotide sequences are also useful as hybridisation probes, as oligomers for PCR, for chromosome and gene mapping, in the recombinant production of protein, or in generating antisense DNA or RNA. The purified polypeptide
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New polynucleotide sequences obtained as hybridization probes, as oligomers mapping, in the recombinant production
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(STAC/)
(DICK/)
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DICKSON M C.
JONES L W.
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hes 0;
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RESULT 131
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Matches 45; Conserv
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27-AUG-1999;
11-JAN-2000;
02-MAY-2000;
09-JUN-2000;
                         length cDNAs defined in the specification. Where a primer set comprises:

(a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide which comprises a 1'-end sequence, where the oligonucleotide comprises a 1'-end sequence and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are are also useful for the sparticularly full-length cDNAs. The primers are are also useful for the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           is useful for generating antibodies specific for it. is one of the 38043 isolated cDNA/SST sequences. Note for this patent did not form part of the printed specobtained in electronic format directly from USPTO at
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Primer sets for synthesizing polynucleotides, length cDNAs defined in the specification, an diagnosis of the abnormality of the proteins
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                                                                                                                                                                                                                                                                                                                                                                                                                         The present invention describes primer sets
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Sugiyama T,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  SEQ
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; 2000JP-00183767.
; 2000JP-00241899.
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99JP-00300253.
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T, Wakamatsu
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A, Nagai K,
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C, Otsuki
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                  The invention relates to a detection reagent capable of detecting one or compared much in the single nucleic acid polymorphisms. The invention also relates to compare single nucleic acid polymorphisms. The invention also relates to compare the sub-region, a trait is linked to one of the human chromosomes or its sub-region, a computer readable medium having stored in it the SNP conclude for detecting at least one SNP given in the specification and isolated nucleic acid molecule for detecting at least one SNP given in the specification comprising at least about 12 contiguous nucleotides, genotyping at least one Contribution in a sample, identifying an individual having or at risk of developing a disorder and a kit comprising at least one container containing the detection reagent. Determining whether a trait is linked to one of the human chromosomes or containing whether the trait is linked to one correctly the detection reagents. Genotyping at least one SNP position given in the specification in a sample comprises contacting the sample with a detection reagent that differentiates between alternative alleger at the least one contacting the sample with a detection reagent that differentiates between alternative
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          New detection reagent capable of detecting 1, 100, 500, 1000 or 5000 or more single nucleic acid polymorphisms, useful in identifying an individual having or at risk of developing type II diabetes or obesity.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        WPI; 2005-511776/52
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10-SEP-2001;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  cic; anorectic; endocrine disease; gastrointestinal disease; disorder; nutritional disorder; single nucleotide polymorphism;
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2001US-00948947.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ; diagnosis; non-insulin dependent diabetes; obesity; anorectic; endocrine disease; gastrointestinal 44000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Liu X,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   601
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position given in
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 English.
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   the specification,
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RESULT 133
AAK63029
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Best Local
                                                                                                                                                                                             26-JUL-2000;
26-JUL-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                                                                                                                                                19-MAY-2000;
07-JUN-2000;
28-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 determining which allele is present at the at least one SNP position. Identifying an individual having or at risk of developing a disorder comprises genotyping at least one SNP given in the specification in a nucleic acid sample from the individual. The disorder is type II diabetes (non-insulin dependent diabetes) or obesity. The detection reagent is useful in identifying an individual having or at risk of developing a disorder, particularly type II diabetes or obesity. This sequence represents a human DNA polymorphic region used in the scope of the invention. Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format from
                                                                                                               14-AUG-2000;
14-AUG-2000;
                                                                                                                                                       14-AUG-2000;
14-AUG-2000;
                                                                                                                                                                                                                                                        11-JUL-2000;
                                                                                                                                                                                                                                                                                                                   07-JUL-2000
                                                                                                                                                                                                                                                                                                                                                             30-JUN-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                           17-MAR-2000;
18-APR-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          02-MAR-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       cytostatic;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Human immune/haematopoietic antigen encoding cDNA SEQ ID NO:8089.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            06-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    AAK63029;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          AAK63029 standard; cDNA; 1664
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Sequence
                                                                                                                                                                                                                                                                                                  11-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           17-JAN-2001;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      2895 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 2939
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               202 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 246
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               at seqdata.uspto.gov/sequence.html.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             45;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       gene
          2000US-0186350P.
2000US-0199874P.
2000US-0199076P.
2000US-0205515P.
2000US-0215135P.
2000US-0216867P.
2000US-0216867P.
2000US-0216867P.
2000US-0217487P.
2000US-0218290P.
2000US-0218290P.
2000US-0218290P.
2000US-022518290P.
2000US-0225213P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           2001WO-US001354
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       therapy; vaccine;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Score 45;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            ВP
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                                                                                                                                                         CC amino acid sequences given in AAM82170 to AAM9121. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) concert and be used to produce the secreted (I), by inserting the culled acids into a host cell and culturing the cell to express the culeic acids into a host cell and culturing the cell to express the concert and treat immune/haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention.
                                                                 Query Match
Best Local S
Matches 45
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17-NOV-2000
17-NOV
                                                                                                                                                       Sequence 1664 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Claim
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2898 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG
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DB; AAM90248.
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                                                                   45; Conserv
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2000US-024920PP.
2000US-024920PP.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
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2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249218P.
2000US-0249218P.
2000US-024924P.
2000US-0249264P.
2000US-0249264P.
2000US-0249269P.
2000US-0249269P.
2000US-0251989P.
2000US-0251866P.
2000US-0251989P.
                                                               1.4%;
llarity 100.0%;
Conservative
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                                                               Score 45; DB
Pred. No. 3.2
0; Mismatches
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B 0;
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                                                                   Indels
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Gaps

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114-AUG-2000
22-AUG-2000
22-AUG-2000
22-AUG-2000
23-AUG-2000
23-AUG-2000
21-SEP-2000
21-SEP-2000
25-SEP-2000
26-SEP-2000
27-SEP-2000
28-SEP-2000
29-SEP-2000
214-SEP-2000
214-SEP-2000
214-SEP-2000
214-SEP-2000
214-SEP-2000
21-SEP-2000

2000US-0225759P.
2000US-022668P.
2000US-022668P.
2000US-02270192P.
2000US-0229343P.
2000US-0229343P.
2000US-0229343P.
2000US-023943P.
2000US-023943P.
2000US-023943P.
2000US-023943P.
2000US-0231043P.
2000US-02312442P.
2000US-02312442P.
2000US-02312443P.
2000US-02312443P.
2000US-02312443P.
2000US-02312443P.
2000US-02312443P.
2000US-0231243P.
2000US-024180P.
2000US-0246474P.
2000US-0246474P.
2000US-0246474P.
2000US-0246525P.
2000US-0246523P.
2000US-0246523P.
2000US-02466528P.
2000US-0246653P.

Human immune/haematopoietic antigen genomic sequence SEQ ID No:24698. W Human; jmmune/haematopoietic jmmune/haematopoietic antigen; cancer; cytostetic; gene therapy; vaccine; metastasis; ds. W Homo sapiems. W HOZOO157182-AZ. HOZOO157182-AZ	333 GGATCACCTG; 134 6 669886 standard; [K69886;
PR 14-SEP-2000; 2000US-0233064P; PR 21-SEP-2000; 2000US-0233064P; PR 21-SEP-2000; 2000US-0234998P; PR 25-SEP-2000; 2000US-0234998P; PR 25-SEP-2000; 2000US-0235836P; PR 27-SEP-2000; 2000US-0235836P; PR 29-SEP-2000; 2000US-0235836P; PR 29-SEP-2000; 2000US-0235359P; PR 29-SEP-2000; 2000US-0235359P; PR 29-SEP-2000; 2000US-0235370P; PR 29-SCT-2000; 2000US-0244950P; PR 29-SCT-2000; 2000US-024478P; PR 20-SCT-2000; 2000US-024478P; PR 20-SCT-2000; 2000US-024478P; PR 20-SCT-2000; 2000US-024479P; PR 20-SCT-2000; 2000US-0244479P; PR 20-SCT-2000; 2000US-02446523P; PR 20-SCT-2000; 2000US-0244523P; PR 20-SCT-2000; 2000US-0244523P; PR 20-SCT-2000; 2000US-0244523P; PR 20-SCT-2000; 2000US-0244231P; PR 20-S	08-SEP-2000; 2000U 08-SEP-2000; 2000U 12-SEP-2000; 2000U 14-SEP-2000; 2000U 14-SEP-2000; 2000U 14-SEP-2000; 2000U 14-SEP-2000; 2000U 14-SEP-2000; 2000U 14-SEP-2000; 2000U

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AAK69885
ID AAK698
XX AAK698
XX AAK698
XX D6-NOV
XX Human
XX Human;
KW Human;
KW Cytost
XX Homo s
XX Homo s
XX HOMO s
XX PN WO2001
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             amino acid sequences given in AAMS2170 to AAMS1921. (1) have cytostatic contivity, and can be used in gene therapy and vaccine production. (1) cytostatic proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (1) expression. For cytostample, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (1) by expressing inactive proteins or to supplement the patients own production of (1). Additionally, (1) polynucleotides may be used to produce the secreted (1), by inserting the culties caids into a host cell and culturing the cell to express the protein. (1) proteins and polynucleotides may be used to prevent, classing and cancer metastases of haematopoietic-derived cells. AAK64703 coancers and cancer metastases of haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                Query Match
Best Local S
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01-DEC-2000
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05-DEC-2000
05-DEC-2000
06-DEC-2000
06-DEC-2000
08-DEC-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                 Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                   Human immune/haematopoietic antigen genomic sequence SEQ ID NO:24697.
                                                                                                                                                                                   06-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence 2219
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     09-AUG-2001.
                                   WO200157182-A2
                                                                    Homo sapiens.
                                                                                                                                                                                                                                                    AAK69885 standard;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    WPI; 2001-483426/52.
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                                                                                                                                                                                                                                                                                                                                                                                                                   Similarity
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2000US-0250391P.
2000US-0251030P.
2000US-0251988P.
2000US-0251479P.
2000US-0251856P.
2000US-0251868P.
2000US-0251869P.
2000US-0251989P.
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2000US-0251989P.
2000US-0251989P.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AAK64702 encode the human immune/haematopoietic antigen (I) sequences given in AAM82170 to AAM91921. (I) have cytostatic
                                                                                                                                                                                                                                                                                                                                                                                                      Conservative
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                                                                                                                                                                                   (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  BP; 633
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                                                                                                                                                                                                                                                    DNA;
                                                                                                                                                                                                                                                                                                                                                                                           1.4%; but
100.0%; Pre
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                                                                                                                                                                                                                                                      2219
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Pred. No.
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3.1e-10;
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     14-AUG-2000
22-AUG-2000
22-AUG-2000
23-AUG-2000
01-SEP-2000
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26-JUL-2000;
26-JUL-2000;
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28-JUN-2000;
30-JUN-2000;
07-JUL-2000;
07-JUL-2000;
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04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAY-2000;
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     2000US-0184664P.
2000US-019875P.
2000US-019875P.
2000US-0199076P.
2000US-0199076P.
2000US-0205115P.
2000US-0215115P.
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2000US-0215115P.
2000US-02116880P.
2000US-0217487P.
2000US-0217487P.
2000US-0225211P.
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2000US-0225211P.
2000US-0225267P.
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2000US-0225768P.
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2000US-023141P.
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2000US-0231243P.
2000US-0231243P.
2000US-0231243P.
2000US-0231414P.
2000US-0231444P.
2000US-02
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17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
01-DEC-2000;
01-DEC-2000;
05-DEC-2000;
05-DEC-2000;
05-DEC-2000;
06-DEC-2000;
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29-SEP-2000
20-CCT-2000
02-CCT-2000
02-CCT-2000
02-CCT-2000
02-CCT-2000
03-CCT-2000
03-NOV-2000
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17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
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17-NOV-2000;
WPI; 2001-483426/52
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                                                                                                                      HUMAN
                                                           Barash SC,
                                                                                                                                                                               2000US-0246609P

2000US-0246611P

2000US-0246611P

2000US-024920PP

2000US-0249211P

2000US-024921P

2000US-024921P

2000US-024926P

2000US-024926P

2000US-024926P

2000US-024926P

2000US-02599P

2000US-02599P

2000US-02599P

2000US-0251988P

2000US-0251866P

2000US-0251869P

2000US-0251989P

2000US-025199PP
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2000US-0239935P

2000US-0241966P

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2000US-0241787P

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2000US-0241809P

2000US-0241878P

2000US-02467476P

2000US-0246476P

2000US-0246478P

2000US-0246478P

2000US-0246528P

2000US-0246528P

2000US-0246528P

2000US-0246532P

2000US-0246532P

2000US-0246532P

2000US-0246532P

2000US-0246532P
                                                                                                                          GENOME SCI INC.
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RESULT 136
AAK86256/c
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         CC amino acid sequences given in AAM82170 to AAM891921. (1) have cytostatic contivity, and can be used in gene therapy and vaccine production. (I) CC proteins and polynucleotides may be used in the prevention, diagnosis and CC treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat disorders associated with decreased CC expression by rectifying mutations or deletions in a patient's genome CC that affect the activity of (I) by expressing inactive proteins or to CC supplement the patients own production of (I). Additionally, (I) CC polynucleotides may be used to produce the secreted (I), by inserting the culcic acids into a host cell and culturing the cell to express the CC protein. (I) proteins and polynucleotides may be used to prevent, CC diagnose and treat immune/haematopoietic-related diseases, especially CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703 CC to AAK87694 represent human immune/haematopoietic antigen genomic CC sequences from the present invention. AAK54950 and AAM82169 CC sequences from the present invention. AAK54950 and AAM82169 CC sequences received cells to protein the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Query Match
             31-JAN-2000

04-FEB-2000

24-FEB-2000

12-MAR-2000

16-MAR-2000

17-MAR-2000

19-MAY-2000

07-JUN-2000

07-JUN-2000

07-JUL-2000

07-JUL-2000

11-JUL-2000

11-JUL-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                  WO200157182-A2
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        cytostatic;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human immune/haematopoietic antigen genomic sequence SEQ ID NO:41068.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      07-NOV-2001
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           AAK86256;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Disclosure; SEQ ID NO 24697; 3071pp + Sequence Listing; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                                                                                                                                                                                                                                                                                                                      17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                             09-AUG-2001.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 represent sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        1966 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTG 2010
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45; Conserv
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2000US-0180628P.
2000US-0186359P.
2000US-0186359P.
2000US-0198123P.
2000US-0205515P.
2000US-0205467P.
2000US-0215135P.
2000US-0211647P.
2000US-0211647P.
2000US-02117486P.
2000US-0217486P.
2000US-0217486P.
2000US-0217486P.
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2000US-0217486P.
2000US-0217486P.
2000US-0217486P.
2000US-0217486P.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          gene therapy; vaccine; metastasis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1.4%; Score 45; DB 4; llarity 100.0%; Pred. No. 3.1e-Conservative 0; Mismatches
                                                                                                                                                                                                                                                                                                                                                        2001WO-US001354.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            559 T; 0 U; 0 Other;
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CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic cactivity, and can be used in gene therapy and vaccine production. (I) CC proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For cexample, they may be used to treat disorders associated with decreased cexpression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) collynucleotides may be used to produce the secreted (I), by inserting the culleic acids into a host cell and culturing the cell to express the collagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metastases of haematopoietic acids encert encert immune/haematopoietic acids encert encert immune/haematopoietic acids encert encert immune/haematopoietic acids encert encert immune/haematopoietic acids encert encert encert encert immune/haematopoietic encert encer
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01.7.NOV-2000
11.7.NOV-2000
01.7.NOV-2000
01.7.NOV-200
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     c acids encoding for preventing,
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2000US-0246477P
2000US-0246523P
2000US-0246532P
2000US-0246532P
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                                                                                                                                                                                                                                                                                                                                                                                                                                    Sequence Listing;
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English.

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and

14-AUG-2000
12-AUG-2000
12-AUG-2000
13-AUG-2000
10-SEP-2000
01-SEP-2000
02-OCT-2000
03-NOV-2000
08-NOV-2000
08-NOV-2000

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2000US-0225214P.
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2000US-0237034P.
2000US-0231243P.
2000US-0231249P.
200

6-JUL-2000; 2000US-022064P. 4-AUG-2000; 2000US-0224518P. 4-AUG-2000; 2000US-0225213P. 4-AUG-2000; 2000US-0225214P. 4-AUG-2000; 2000US-0225266P. 4-AUG-2000; 2000US-0225268P. 4-AUG-2000; 2000US-0225268P. 4-AUG-2000; 2000US-0225757P. 4-AUG-2000; 2000US-0225759P. 4-AUG-2000; 2000US-0225759P. 4-AUG-2000; 2000US-0225759P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-0225759P. 4-AUG-2000; 2000US-0225759P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-022679P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-022668P. 4-AUG-2000; 2000US-022668P. PR 4-AUG-2000; 2000US-022668P. PR 2-AUG-2000; 2000US-022688P. PR 2-AUG-2000; 2000US-022688P. PR 2-AUG-2000; 2000US-022688P. PR 2-AUG-2000; 2000US-022688P. PR 2-AUG-2000; 2000US-02268P. PR 2-AUG-2000; 2000US-022934P. PR PR 1-SEP-2000; 2000US-022934P. PR PR PR PR PR PR PR PR PR	31-JAN-2000; 2000US-0179065P. 04-FEB-2000; 2000US-0184664P. 24-FEB-2000; 2000US-01846350P. 16-MAR-2000; 2000US-018974P. 17-MAR-2000; 2000US-0198123P. 18-AFR-2000; 2000US-0198123P. 19-MAY-2000; 2000US-0205515P. 07-JUN-2000; 2000US-020467P. 28-JUN-2000; 2000US-0214866P. 30-JUN-2000; 2000US-0215135P. 07-JUL-2000; 2000US-0216647P. 07-JUL-2000; 2000US-0216647P. 11-JUL-2000; 2000US-0217487P. 11-JUL-2000; 2000US-0217487P. 11-JUL-2000; 2000US-0217495P. 11-JUL-2000; 2000US-0217495P. 11-JUL-2000; 2000US-0217495P. 11-JUL-2000; 2000US-0218290P. 12-JUL-2000; 2000US-0218290P.	## Homo sapiens. PR ## W0200157182-A2. PR ## 09-AUG-2001. PR P	AAK86257; PRAAK86257; PRAAK86257; PRAAK86257; PRAAK86257; PRO7-NOV-2001 (first entry) PRHuman immune/haematopoietic antigen genomic sequence SEQ ID NO:41069. PRHuman; immune; haematopoietic; immune/haematopoietic antigen; cancer; PRCytostatic; gene therapy; vaccine; metastasis; ds. PRAK86257; PR	Sequence 2219 BP; 559 A; 519 C; 508 G; 633 T; 0 U; 0 Other; PR Sequence 2219 BP; 559 A; 519 C; 508 G; 633 T; 0 U; 0 Other; PR Query Match 1.4%; Score 45; DB 4; Length 2219; Best Local Similarity 100.0%; Pred. No. 3.1e-10; PR Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0; PR Matches 45; TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGGACCAGCCTG 2932
08-NOV-2000 17-NOV-2000	02-0CT-2000; 02-0CT-2000; 02-0CT-2000; 02-0CT-2000; 13-0CT-2000; 13-0CT-2000; 20-0CT-2000;	27-SEP-2000; 27-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000; 29-SEP-2000;	14-SEP-2000; 14-SEP-2000; 14-SEP-2000; 14-SEP-2000; 14-SEP-2000; 21-SEP-2000; 21-SEP-2000; 21-SEP-2000; 25-SEP-2000; 25-SEP-2000;	
2000US -0246476P. 2000US -0246478P. 2000US -0246524P. 2000US -0246524P. 2000US -0246526P. 2000US -0246527P. 2000US -024652P. 2000US -024652P. 2000US -024652P. 2000US -024652P. 2000US -024652P. 2000US -024651P. 2000US -024651P. 2000US -024661P. 2000US -024661P. 2000US -024621P. 2000US -024921P.				

17-NOV-2000; 17-NOV-2000; 17-NOV-2000; 17-NOV-2000;

17-NOV-2000;

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                                                                                                                                                                                                                                                                                                        CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic cc activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For c example, they may be used to treat disporders associated with decreased cc expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the cc nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, cd diagnose and treat immune/haematopoietic-derived cells. AAK64703 c cancers and cancer metastases of haematopoietic antigen genomic cancers from the present inwune/haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention
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17-NOV-2000;
17-NOV-2000;
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01-DEC-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                04-NOV-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 41069; 3071pp + Sequence Listing; English
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                                               ADR07058
                                                                            ADR07058 standard;
                                                                                                                                                                                                                                                                                  Sequence 2219 BP; 559 A; 519 C; 508 G; 633 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               WPI; 2001-483426/52.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             (HUMA-) HUMAN GENOME SCI INC
                                                                                                                                                                                        2888 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTG 2932
                                                                                                                                                          254
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                                                                                                                                                                                                                     45;
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2000US-0249218P.
2000US-0249218P.
2000US-024924P.
2000US-0249248P.
2000US-0249264P.
2000US-0249265P.
2000US-0249299P.
2000US-0249299P.
2000US-0249299P.
2000US-0259300P.
2000US-0251930P.
2000US-025193P.
2000US-025198P.
2000US-0251866P.
2000US-0251866P.
2000US-0251989P.
2000US-0259678P.
                                                                                                                                                                                                                    Conservative
                (first entry)
                                                                            CDNA; 2491
                                                                                                                                                                                                                                    1.4%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Ruben
                                                                                                                                                                                                                                    Score 45; ; Pred. No.
                                                                                                                                                                                                                    0
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                                                                            ВP
                                                                                                                                                                                                                     Mismatches
                                                                                                                                                                                                                                    3.1e-10;
                                                                                                                                                                                                                                                   DB 4;
                                                                                                                                                                                                                     0
                                                                                                                                                                                                                                                 Length 2219;
                                                                                                                                                                                                                     Indels
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                                                                                                                                                                                                                    Gaps
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8

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Human cDNA sequence SEQ ID NO:18165

26-JUN-2001 AAH18230

(first entry)

AAH18230 standard; cDNA;

3977

ВP

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RESULT 139
AAH18230
ID AAH182
XX
AC AAH182
XX
DT 26-JUN
XX
DB Human
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Query Match
Best Local (
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                to and modulate expression of the CDNA molecules. As such, these molecules are useful for diagnostic markers or therapeutic targets for the various diseases or morbid states. In particular, they are useful in gene therapy for treating osteoporosis, neurological disease, Alzheimer's disease, Parkinson's disease, dementia, short memory and various cancers, as well as for maintaining equilibrium of sense or motor function, and for treating emotional reaction, fear response and panic. Accordingly, they exhibit osteopathic, neuroprotective, nootropic, antiparkinsonian, cytostatic and tranquiliser activities. This polymucleotide is a full length human cDNA sequence of the invention. NOTE: This sequence is not given in the sequence listing of the specification but can be obtained on CD-ROM from the European Patent Office, Vienna Sub-office.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  This invention relates to novel, isolated full length human cDNA molecules and the encoded proteins thereof. Specifically, it refers to molecules and the encoded proteins thereof. Specifically, it refers to molecules and obtained by an oligo-capping method, where none of these clones are identical to any known human mRNAs. The present invention describes an immunoassay to identify agonists and antagonists, as well antibodies, antisense molecules and siRNAs that can all be used to bind antibodies, antisense molecules and siRNAs that can all be used to bind
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 1; SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    New 1995 cDNA, useful for treating osteoporosis, neurological diseases, Alzheimer's diseases, Parkinson's diseases, dementia and various cancers.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           WPI; 2004-583265/57.
P-PSDB; ADR09014.
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09-MAY-2003; 2003JP-00131452.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            18-AUG-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  EP1447413-A2
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Parkinson's disease; dementia; short memory; cancer; sense or motor function; emotional reaction; fear response; osteopathic; neuroprotective; nootropic; antiparkinsonian;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Full length human cDNA useful for treating neurological disease Seq 564.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Wakamateu
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Isogai T,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                12-FEB-2004; 2004EP-00003145
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            osteoporosis; neurological
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      (REAS-) RES ASSOC BIOTECHNOLOGY
                                                                                                                                                                                                                                                                                                                                                        2447
                                                                                                                                                                                                                                                                                                                                                                                      3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ss; human; oligo-capping method; diagnostic
                                                                                                                                                                                                                                                                                                                                                                                                                                                              l Similarity
45; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   2491 BP; 509 A; 639 C; 794 G; 549 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                     GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 2491
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Yamamoto J,
A, Ishii S,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              NO 564; 2686pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            1.4%;
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Nagai K, Irie
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            disease; Alzheimer's disease;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Score 45;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Isono Y, Sugiyama T, e R;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  DB 13;
3.1e-10
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               .1e-10;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Indels
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RESULT 140
AAK69446/c
ID AAK694
XX
AC AAK694
XX
DT 06-NOV
XX
DH Human
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                                                                                                                                                                                                                                                                                                         length cDNAs defined in the specification. Where a primer set comprises:

((a) an oligo-dr primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 complementary strand of a polynucleotide which comprises one of the 5602 coligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end complementary to a polynucleotide which comprises a 3'-end sequence complementary to a golynucleotide which comprises a 3'-end sequence complementary to a coligonucleotide which comprises a 3'-end sequence, where the coligonucleotide which comprises at least 15 nucleotides and the combination of the 5'-end sequence) are sets can be used in antisense therapy and in the specification. The primers are useful for synthesising polynucleotides, comparisons the primers are also useful for the sparticularly full-length cDNAs. The primers are also useful for the comprise comparisons the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs assity without any specialised methods. AAH03166 to AAH13628 and CC DNAs easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods. AAH03165 to AAH13628 and CC DNAS easily without any specialised methods.
                                                                                                                                                                                                                              Query Match
Best Local (
                                                                                                                                                                                                                Matches
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27-AUG-1999; 99JP-00300253.
11-JAN-2000; 2000JP-00118776.
02-MAY-2000; 2000JP-00183767.
09-JUN-2000; 2000JP-00241899.
 Human immune/haematopoietic antigen genomic sequence SEQ ID NO:24258
                             06-NOV-2001
                                                         AAK69446;
                                                                                    AAK69446
                                                                                                                                                                                                                                                                        Sequence 3977
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Claim
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Primer sets for synthesizing polynucleotides, particularly the 5602 full-
length cDNAs defined in the specification, and for the detection and/or
diagnosis of the abnormality of the proteins encoded by the full-length
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     WPI; 2001-318749/34.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        28-JUL-2000; 2000EP-00116126
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    07-FEB-2001.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Human; primer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          (HELI-) HELIX RES INST.
                                                                                                                                                                                                                               Local Similarity
                                                                                                                                                         3931
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       present invention
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   8
                                                                                                                                                                                                                                                                                                     invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Isogai T,
                                                                                                                                                                     GTGCCACTGCACTCCAGCCTGGGCAACAGAGACAGACTCTGTCTC 3122
                                                                                  standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                SEQ ID NO 18165; 2537pp +
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                                                                                                                                                                                                                Conservative
                            (first entry)
                                                                                                                                                                                                                                                                       BP; 1063 A; 888 C; 987 G; 1039 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  detection; diagnosis; antisense therapy; gene therapy; ss
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               'n
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Nishikawa
                                                                                    DNA;
                                                                                                                                                                                                                              100.0%;
                                                                                                                                                                                                                              1.4%; Score 45;
100.0%; Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     describes primer sets for synthesising 5602 full-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Wakamatsu A,
                                                                                                                                                                                                                0; Mismatches
                                                                                    ΒP
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                                                                                                                                                                                                                           DB 4; L
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C, Otsuki
                                                                                                                                                                                                                <u>,</u>
                                                                                                                                                                                                                                           Length 3977;
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Human; immune; haematopoietic; immune/haematopoietic antigen; cytostatic; gene therapy; vaccine; metastasis; ds.
                                              cancer;
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Homo sapiens.

WO200157182-A2

2001WO-US001354

30-JUN-2000; 07-JUL-2000; 07-JUL-2000; 28-JUN-2000; 2000US-0180628P 2000US-0184664P 2000US-018974P 2000US-0198123P 2000US-0214886P 2000US-021680P 2000US-021680P 2000US-021680P 2000US-0217487P 2000US-0217487P 2000US-0217487P 2000US-0218290P 2000US-022963P 2000US-022963P 2000US-0225213P 2000US-0235213P 2000US-023531414P 2000US-0231414P 2000US-0231414P 2000US-0231249P 2000US-0231249P 2000US-0231249P 2000US-0231249P 2000US-0231249P 2000US-0232399P 2000US-0232399P 2000US-02332401P 2000US-02332401P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P 2000US-0233239P

11-JUL-2000; 11-JUL-2000; 14-JUL-2000; 26-JUL-2000;

14-AUG-2000; 14-AUG-2000; 14-AUG-2000;

14-AUG-2000; 14-AUG-2000; 14-AUG-2000; 14-AUG-2000; 14-AUG-2000; 14-AUG-2000;

14-AUG-2000;
18-AUG-2000;
22-AUG-2000;
22-AUG-2000;
23-AUG-2000;
30-AUG-2000;
01-SEP-2000;
01-SEP-2000;
01-SEP-2000;
05-SEP-2000;
06-SEP-2000;
06-SEP-2000;
08-SEP-2000;
08-SEP-2000;
08-SEP-2000;
14-SEP-2000;

2000US-0234274P.
2000US-0234997P.
2000US-023499P.
2000US-0235834P.
2000US-0235834P.
2000US-02358367P.
2000US-0236367P.
2000US-0236369P.
2000US-0236369P.
2000US-0236369P.
2000US-0236369P.
2000US-0236369P.

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RESULT 141
AAL03041/c
ID AAL030
XX AAL030
XX AAL030
XX Human
XX Human
XX Human
XX Homo 8
XX Homo 9
XX Homo 9
XX Homo 9
XX Homo 10-AUG
XX Homo 9
XX Homo 10-AUG
XX Homo 11-AUG
PR 11-AUG
PR
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) antino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) croteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) concleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynuclootides may be used to prevent. (C diagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metasteses of haematopoietic artigen genomic concers and treat immune/haematopoietic artigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Query Match
Best Local Similarity
31-JAN-2000;
04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAY-2000;
07-JUN-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            AAL03041 standard; DNA; 6565
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 4513 BP; 897 A; 1168 C; 1278 G; 1170 T; 0 U; 0 Other;
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05-JAN-2001; 2001US-0259678P.
                                                                                                                                                                                                                                                                            17-JAN-2001;
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                                                                                                                                                                                                                                                                                                                                 02-AUG-2001.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Human reproductive system related antigen DNA SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    21-NOV-2001
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               gene therapy;
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  2000US-0179065P.
2000US-0180628P.
2000US-0184664P.
2000US-0186350P.
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2000US-0190076P.
2000US-0198123P.
2000US-0205515P.
2000US-0209467P.
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2000US-0236802P.
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2000US-0249268P.
2000US-0251868P.
2000US-0251868P.
2000US-0251980P.

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11-NO
                                   The present invention provides the protein and coding sequences of number of human reproductive system related antigens. These can hin the prevention and treatment of reproductive system disorders, including cancer. The present sequence is a genomic sequence encoprotein of the invention
                                                                                                                                                                                                        Isolated
used in p
Sequence 6565
                                                                                                                                                                  Disclosure;
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                                                                                                                                                                                                        preventing,
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English.

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30-JUN-2000
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2000US-021486P 2000US-021647P 2000US-021664PP 2000US-02164PP 2000US-02164PP 2000US-02164PP 2000US-02164PP 2000US-0217496P 2000US-0225451PP 2000US-022521PP 2000US-022521PP 2000US-022521PP 2000US-022526PP 2000US-022526PP 2000US-022526PP 2000US-022526PP 2000US-022575PP 2000US-022575PP 2000US-022575PP 2000US-022575PP 2000US-022575PP 2000US-0227182PP 2000US-0227182PP 2000US-0231411PP 2000US-023141PP 2000US-0231411PP 2000US-023141PP 2000US

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06-SEP-2000 08-SEP-2000 08-SEP-2000 08-SEP-2000 08-SEP-2000 08-SEP-2000 08-SEP-2000 11-SEP-2000 11-SEP-2000 11-SEP-2000 11-SEP-2000 11-SEP-2000 11-SEP-2000 11-SEP-2000 21-SEP-2000 21-SEP-2000 22-SEP-2000 23-SEP-2000 25-SEP-2000 25-SEP-2000 25-SEP-2000 26-SEP-2000 27-SEP-2000 29-SEP-2000 29-SEP-2000 20-OCT-2000 20-OCT-2000 20-OCT-2000 02-OCT-2000 02-OCT-2000 02-OCT-2000 02-OCT-2000 02-OCT-2000 03-NOV-2000 03-NOV-2000 04-NOV-2000 06-NOV-2000 06-NOV-2000 06-NOV-2000 07-NOV-2000 08-NOV-2000 01-NOV-2000	06-SEP-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      The present invention provides the protein and coding sequences of a number of human reproductive system related antigens. These can be used in the prevention and treatment of reproductive system disorders, including cancer. The present sequence is a genomic sequence encoding a protein of the invention
                    17-JAN-2001; 2001WO-US001329
                                                             02-AUG-2001.
                                                                                                                                                                         reproductive system disorder; urinary system disorder; gene therapy; cardiovascular disorder; respiratory disorder; neurological disorder; gastrointestinal disease; infection; cytostatic; gene; ds.
                                                                                                                                                                                                                                  Human; testicular antigen; testes; cancer; metastasis; immune disorder;
                                                                                                                                                                                                                                                                      Human testicular antigen encoding
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Isolated nucleic acid molecule encoding a reproductive system antigen used in preventing, treating or ameliorating a medical condition.
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                                                                                                                                                                                                                                                                                                                                                                                           standard;
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ilarity 100.0%;
Conservative (
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2000US-0249245P.
2000US-0249264P.
2000US-0249265P.
2000US-0249299P.
2000US-0249299P.
2000US-0259100P.
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2000US-02511989P.
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                                                                                                                                                                                                                                                                                                                                                                                           DNA;
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                                                                                                                                                                                                                                                                                                                                                                                             6565
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3e-10;
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2000US-0179065P

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English

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08-NOV-2000
09-NOV-2000
017-NOV-2000
117-NOV-2000
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117-NOV-2000
117-NOV-2000
01-DEC-2000
01-DEC-2000
01-DEC-2000
01-DEC-2000
08-DEC-2000
                Nucleic acids encoding 973 human testicular antigen polypeptides, useful for preventing, diagnosing and/or treating testicular cancer.
                                                                     WPI; 2001-483232/52.
                                                                                                                                          HUMAN
                                                                                                        Barash
                                                                                                                                                                                         2000US-0237037P.
2000US-0237038P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human; testicular antigen; testes; cancer; metastasis; immune disorder; resproductive system disorder; urinary system disorder; gene therapy; cardiovascular disorder; respiratory disorder; neurological disorder; gastrointestinal disease; infection; cytostatic; gene; ds.
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45; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            standard;
2000US-0179065P.
2000US-0184664P.
2000US-0184654P.
2000US-0184654P.
2000US-0184657P.
2000US-0199076P.
2000US-0214886P.
2000US-0216647P.
2000US-0216647P.
2000US-0217486P.
2000US-0217486P.
2000US-0217496P.
2000US-0217496P.
2000US-0217496P.
2000US-0217496P.
2000US-025751P.
2000US-0255266P.
2000US-025547P.
2000US-0225758P.
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Pred. No. 3e-
0; Mismatches
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ВP

DB 4; 3e-10;

Length 6565, Indels

<u>.</u>.

0

Gaps

0

5096 3122

DNA fragment SEQ

ID NO:

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RESULT 145
AAX23521
ID AAX235
XX
AC AAX235
XX
AC AAX235
XX
DT 17-JUN
XX
                                                                                                                                                                                                         밁
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                                                                                                                                                                                                                                                                                                                                                                           Query Match 1.4%; So
Best Local Similarity 100.0%; F
Matches 45; Conservative 0;
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08-NOV-2000
17-NOV-2000
17-NOV
                                                                                                                                                                                                                                                                                                                                                                                                                  protein
                                                                                                                                                                                                                                                                                                                                                                                                                                       gastrointestinal disorders, infecti especially testicular cancers. The
                                                                                                                                                                                                                                                                                                                                                                                                                                                                            The present invention provides the protein and coding sequences of 973 human testicular antigens, and fragments of their genomic sequences. The sequences can be used in the treatment of cardiovascular, urinary system, reproductive system, immune, respiratory, neurological and
                   17-JUN-1999
                                                            AAX23521;
                                                                                                   AAX23521
                                                                                                                                                                                                                                                                                                                                                                        Sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Disclosure;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         (HUMA-) HUMAN GENOME
                                                                                                                                                                                                           5140
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           leic acids encoding 973 human testicular antigen polypeptides, useful preventing, diagnosing and/or treating testicular cancer.
                                                                                                                                                                                                                                               3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         2001-483232/52.
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                                                                                                                                                                                                                                                                                                                                                                                                               fragment
                                                                                                     standard;
                                                                                                                                                                                                         6565 BP; 1349
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2000US-0246613P.
2000US-024920P.
2000US-024920P.
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2000US-0249211P.
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2000US-0249211P.
2000US-0249211P.
2000US-024921P.
2000US-024921P.
2000US-024921P.
2000US-024921P.
2000US-024924P.
2000US-024924P.
2000US-0249264P.
2000US-0249264P.
2000US-0249269P.
2000US-025939P.
2000US-025939P.
2000US-0251866P.
2000US-0251866P.
2000US-0251869P.
2000US-0251869P.
2000US-0251989P.
                     (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                  of the invention
                                                                                                   DNA; 16595
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                                                                                                                                                                                                                                                                                                                                                                                                                                     infections, and rs. The present a
                                                                                                                                                                                                                                                                                      Score 45; DB 4; Le
; Pred. No. 3e-10;
0; Mismatches 0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                       sequence is a DNA encoding
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                                                                                                                                                                                                           5096
                                                                                                                                                                                                                                                                                      0,
                                                                                                                                                                                                                                                                                      Gaps
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14 AUG-2000
22-AUG-2000
22-AUG-2000
22-AUG-2000
23-AUG-2000
23-AUG-2000
21-SEP-2000
01-SEP-2000
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05-SEP-2000
06-SEP-2000
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01-SEP-2000
01-SEP-2000
02-CCT-2000
03-NCV-2000
03-NCV-2000
08-NCV-2000

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2000US-022934P.
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2000US-022934P.
2000US-023934P.
2000US-023934P.
2000US-023124P.
2000US-023141P.
2000US-02411P.
2000US-02411P.
2000US-024647P.
2000US-024647P.
2000US-024647P.
2000US-024652P.
2000US-024652P.
2000US-024652P.
2000US-024663P.

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RESULT 146
AAS36670
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12252222222222
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                                                                                                                                                                                                             S
                                                                                                                                                                                                                                                                                                             CC (Amp). This protein is used to produce recombinant Amp and can be used CC (Amp). This protein is used to produce recombinant Amp and can be used CC for gene therapy for treating Amp-deficiency conditions. Its fragments CC are used as primers and probes to identify patients with homozygous and heterozygous Amp deficiency, including prenatal diagnosis (patients CC defective in Amp are at risk of developing angioedema if treated with CC angiotensin-converting enzyme inhibitors), also as antisense inhibitors CC in cases of excessive Amp expressing sequences in other animals and to generate transgenic animals, and comparisons of genomic sequences are used to detect mutations. Amp inhibitors are potentially useful as antihypertensive agents and to prevent or treat arterial (re)stensis or atherosclerosis. The structure of Amp is used to design synthetic substrates, e.g. for use in Amp assays. Amp, which hydrolyzes N-terminal indo bonds, can be used to degrade industrial protein feeds to free formulations used to treat malabsorption syndrome and for studying its biological role. Antibodies against Amp are used in immunohistochemical
                                                                                                                                                                                                                                    Query Match
Best Local S
Matches 45
         Cardiovascular system antigen; human; mouse; rabbit; goat; horse; cat; chicken; sheep; immunosuppressive; antiarthritic; vasotropic; dog; antirheumatic; antiproliferative; cytostatic; cardiant; neuroprotective; cerebroprotective; nootropic; antibacterial; virucide; fungicide; cancer;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            02-SEP-1997;
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                                                                                       17-DEC-2001
                                                                                                                                   AAS36670 standard; DNA; 17581
                                                                                                                                                                                                                                                                               Sequence 16595
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ophthalmological;
                                                                                                             AAS36670
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                                                                                                                                                                                          4301
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                                                                  cardiovascular
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     acid
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                                                                                                                                                                                                                                                 Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        diagnosis; angioedema; antihypertensive agent; atherosclero
stenosis; industrial protein feed; malabsorption syndrome;
ceous waste degradation; additive; immunohistochemistry; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Sprinkle TJC,
                                                                                                                                                                                                                                                                                                     role.
                                                                                                                                                                                                                                      Conservative
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                                                                                       (first entry)
                                                                                                                                                                                                                                                                                BP; 4429 A; 4145 C;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          human;
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vulnerary; gene therapy; autoimmune disease; neoplasm;
                                                                 system
                                                                                                                                                                                                                                                100.0%;
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; Pred. No.
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                                                                 antigen genomic DNA SEQ ID
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feed; malabsorption syndrome;
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hyperproliferative disorder; breast; liver; cardiovascular disorder; d cerebrovascular disorder; nervous system disorder; bacterial infection fungal infection; ocular disorder; bacterial disorder gastrointestinal disorder; renal disorder; respiratory disorder; wound healing; skin aging; organ transplantation; tissue regeneration; anti-infertility. disorder;

02-AUG-2001

17-JAN-2001; 2001WO-US001340

16-MAR-2000
18-MAR-2000
18-MAR-2000
18-MAR-2000
19-MAR-2000
20-JUN-2000
20-JUN-2000
11-JUL-2000
11-JUC-2000
11-JUC 2000US-0189874P.
2000US-0198123P.
2000US-0205515P.
2000US-021486P.
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2000US-02293

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2000US-0232080P

; 2000US-0232399P. ; 2000US-0232400P. ; 2000US-0232401P. 2000US-0233063P.

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2000US-0234274P.
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2000US-0246477P.
2000US-0246478P.
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2000US-0240960P.
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2000US-0239935P.
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08-DEC-2000;
08-DEC-2000;
08-DEC-2000;
11-DEC-2000;
05-JAN-2001;
                            cardiac arrest, cerebrovascular disorders such as cerebral ischaemia, nervous system disorders such as Alzheimer's disease, infections caused by bacteria, viruses and fungi, ocular disorders such as corneal infection, endocrine disorders such as premature labour and infertility, gastrointestinal disorders such as crohn's disease, renal disorders such as glomerulomephritis and respiratory disorders such as asthma and pleurisy. The polypeptides can also be used to aid wound healing, to prevent skin aging due to sumburn, to maintain organs before transplantation, to regenerate tissues and in chemotaxis. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO
                                                                                                                                                                                                                                                                  disorders in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. A pathological condition can be determined by detecting the presence or absence of a mutation in a cardiovascular system antigen polynucleotide. The treatable disorders include autoimmune diseases such as rheumatoid arthritis, hyperproliferative disorders such as neoplasms of the breast or liver, cardiovascular disorders such as cardiac arrest, cerebrovascular disorders such as cerebral ischaemia.
                                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequences AAS35741-AAS36942 represent genomic DNA molecules, which encounthe cardiovascular system antigen polypeptides of the invention. Cardiovascular system antigens and their associated polynucleotides are
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   New cardiovascular system related polynucleotides and polypeptides, useful for diagnosing, treating and/or preventing disorders of the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              cardiovascular system.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  (HUMA-) HUMAN GENOME SCI INC.
              ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                               in the diagnosis, treatment and prevention of various types
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 SEQ ID NO 2170; 674pp; English.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ; 2000US-0251869P.
; 2000US-0251989P.
; 2000US-0251990P.
; 2000US-0254097P.
; 2001US-0259678P.
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encode

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                  S
                                 Query Match
Best Local Similarity
Matches 45; Conser
4841
                    2898 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 2942
 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 4885
                                          Conservative
                                    1.4%; 500
100.0%; Pr
                                       Score 45; DB; Pred. No. 2.9
0; Mismatches
                                      DB 4;
3. 2.9e-10;
0;
                                                             Length 17581;
                                           Indels
                                          0
                                           Gaps
                                           0
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AAK83984 standard; DNA; 17581 BP

07-NOV-2001 (first entry)

Human immune/haematopoietic antigen genomic sequence SEQ ID NO:38796.

RESULT 147
AAK83984
ID AAK839
XX AAK839
XX O7-NOV
XX O7-NOV
XX Human
XX Human
XX Human
XX Horost
XX O2001
XX O2001
XX Horost
X Human; immune; haematopoietic;
cytostatic; gene therapy; vacci vaccine; immune/haematopoietic antigen; cancer; ine; metastasis; ds.

Homo sapiens.

WO200157182-A2

17-JAN-2001; 2001WO-US001354.

31-JAN-2000; 2000US-0179065P

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04.FBB-2000
02.4PBB-2000
02.4PBB-2000
02.4PBB-2000
01.4PBB-2000
11.7MAR-2000
01.7MAR-2000
01.7MA
2000US-0186628

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2000US-0186359

2000US-0198139

2000US-0295159

2000US-02151359

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2000US-02151359

2000US-02168809

2000US-0218290

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2000US-0225217496

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2000US-023342399

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2000US-02334239

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2000US-0236369

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    02-OCT-2000

02-OCT-2000

13-OCT-2000

20-OCT-2000

20-OCT-2000

20-OCT-2000

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20-OCT-2000

20-OCT-2000

20-OCT-2000

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01-NOV-2000

01-NO
                                           Nucleic
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  Disclosure; SEQ ID NO 38796; 3071pp + Sequence Listing; English.
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                                         c acids encoding for preventing,
                                                                                                                                                                                                     HUMAN
                                                                                                                                                         Barash
                                                                                                                                                                                                                                            2000US-0237038P.
2000US-0237039P.
2000US-0239935P.
2000US-0241966P.
2000US-0241986P.
2000US-0241809P.
2000US-02418109P.
2000US-0246477P.
2000US-0246477P.
2000US-0246477P.
2000US-0246477P.
2000US-0246478P.
2000US-0246524P.
2000US-0246524P.
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2000US-0246524P.
2000US-0246524P.
2000US-0246521P.
2000US-024651P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-0249211P.
2000US-024921P.
2000US-025921P.
2000US-02592P.
2000US-0
                                                                                                                                                                                                     GENOME
                                                                                                                                                         SC,
                                                                                                                                                                                                       SCI
                                         human immune/hematopoietic diagnosing and/or treating
                                                                                                                                                            Ruben
                                                                                                                                                                                                     INC.
                                                                                                                                                            MS.
                                             antigen polypeptides, cancers and metastasis.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    RESULT 148
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       CC AAKS4951 to AAK64702 encode the human immune/haematopoietic antigen (I) CC amino acid sequences given in AAW82170 to AAW91921. (I) have cytostatic CC activity, and can be used in gene therapy and vaccine production. (I) CC proteins and polymucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat disorders associated with decreased CC expression by rectifying mutations or deletions in a patient's genome CC that affect the activity of (I) by expressing inactive proteins or to Supplement the patients own production of (I). Additionally, (I) CC supplement the patients own production of (I). Additionally, (I) CC polynucleotides may be used to proved the secreted (I), by inserting the CC nucleic acids into a host cell and culturing the cell to express the CC diagnose and treat immune/haematopoietic-derived cells. AAK64703 CC cancers and cancer metastases of haematopoietic antigen genomic CC sequences from the present invention. AAK54942 to AAK8769 and AAW82169
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Query Match
Best Local S
Matches 45
 16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAY-2000;
07-JUN-2000;
28-JUN-2000;
30-JUN-2000;
07-JUL-2000;
07-JUL-2000;
11-JUL-2000;
11-JUL-2000;
                                                                                                                                                                      31-JAN-2000;
04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
                                                                                                                                                                                                                                                                                                                                                                                                            Human; cardiovascular system related polypeptide; cancer; proliferative disorder; foetal abnormality; developmental abnormality; haematopoietic disorder; AIDS; autoimmune disease; rheumatoid arthritis; inflammation; allergy; neurological disorder; Alzheimer's disease; Parkinson's disease; cognitive disorder; schizophrenia; asthma; skin disorder; psoriasis; sepsis; diabetes; atherosclerosis; cardiovascular disorder; angiogenic disorder; kidney disorder;
                                                                                                                                                                                                                                                     07-MAR-2002;
                                                                                                                                                                                                                                                                                    27-MAR-2003.
                                                                                                                                                                                                                                                                                                                  US2003059908-A1
                                                                                                                                                                                                                                                                                                                                                 Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                gastrointestinal disorder;
endocrine disorder; gene; d
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human cardiovascular system related genomic DNA #930.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            29-JAN-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        ADE47364;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ADE47364 standard; DNA; 17581
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Sequence 17581 BP; 4762 A; 3663 C; 4018 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              represent sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  4841 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 4885
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   2898
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45; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 2942
2000US-0184664P
2000US-0184664P
2000US-0186350P
2000US-0189874P
2000US-0199874P
2000US-0198123P
2000US-0205515P
2000US-020567P
2000US-02148467P
2000US-0216647P
2000US-021684P
2000US-0217486P
2000US-0217486P
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                                                                                                                                                                                                                                                     2002US-00091504.
                                                                                                                                                                                                                    2000US-0179065P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         (first entry)
                                                                                                                                                                                                                                                                                                                                                                              gene; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              1.4%;
                                                                                                                                                                                                                                                                                                                                                                                              pregnancy-related disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               0,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Score 45;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       BP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DB 4; Le
. 2.9e-10;
thes 0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Length 17581;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               0;
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   14-SBP-2000
14-SEP-2000
14-SEP-2000
21-SEP-2000
21-SEP-2000
25-SEP-2000
27-SEP-2000
27-SEP-2000
29-SEP-2000
21-OCT-2000
02-OCT-2000
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08-SEP-2000;
08-SEP-2000;
08-SEP-2000;
08-SEP-2000;
08-SEP-2000;
08-SEP-2000;
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26-JUL-2000

26-JUL-2000

14-AUG-2000

12-AUG-2000

12-AUG-2000

22-AUG-2000

22-AUG-2000

23-AUG-2000

23-AUG-2000

21-SEP-2000

01-SEP-2000

01-SEP-2000
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14-SEP-2000;
14-SEP-2000;
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              2000US-023423P.
2000US-023499PP.
2000US-023499PP.
2000US-0235834P.
2000US-0235834P.
2000US-0235834P.
2000US-023636PP.
2000US-023636PP.
2000US-023637PP.
2000US-023637PP.
2000US-023637PP.
2000US-023637PP.
2000US-023637PP.
2000US-023703PP.
2000US-0241786PP.
2000US-0241786PP.
2000US-0241809P.
2000US-0241809P.
2000US-0241809P.
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2000US-022451BP

2000US-0225213P

2000US-022526P

2000US-022526P

2000US-022526P

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2000US-022575P

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2000US-0226868P

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2000US-023934P

2000US-023939P

2000US-0231243P

2000US-0231413P

2000US-02314149P

2000US-02314149P

2000US-023140P

2000US-023240P

2000US-023340P

2000US-023340P
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08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
17-NOV-2000
17-NOV
                                                   The invention relates to human cardiovascular system related polypeptides and the polynucleotides encoding them. The polypeptides, polynucleotides and antibodies to the polypeptides are useful for diagnosing a pathological condition or a susceptibility to a pathological condition, for preventing, treating, or ameliorating a medical condition, such as cancer of cardiovascular system tissues, proliferative disorders, foetal and developmental abnormalities, haematopoietic disorders, diseases of the immune system, AIDS, autoimmune diseases (e.g., rheumatoid the immune system, AIDS, autoimmune diseases (e.g., rheumatoid
                                                                                                                                                                                                                                                                              New cardiovascular system related polynucleotides and polypeptides, useful for preventing, treating, or ameliorating a medical condition, such as cancer of cardiovascular tissues and cancer metastases.
            arthritis), inflammation, allergi
Alzheimer's disease, Parkinson's
                                                                                                                                                                                                                                                                                                                                                                                                      Rosen
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   -NOV-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           -NOV-2000;
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                                                                                                                                                                                                                                                                                                                                                                 2003-743766/70.
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                                                                                                                                                                                                                                                                                                                                                                                                                                              HUMAN
                                                                                                                                                                                                                                          SEQ
                                                                                                                                                                                                                                                                                                                                                                                                    Ruben SM,
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2000US-0246475P.
2000US-0246478P.
2000US-0246524P.
2000US-0246528P.
2000US-0246528P.
2000US-0246510P.
2000US-0246511P.
2000US-0246511P.
2000US-0246511P.
2000US-0246511P.
2000US-0246511P.
2000US-0249211P.
2000US-0249218P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0249219P.
2000US-0251988P.
2000US-0251868P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
2000US-0251989P.
                               inflammation, allergies, neurological disorders (e.g.,
                                                                                                                                                                                                                                             ä
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     2001US-00764869
                                                                                                                                                                                                                                                                                                                                                                                                                                              GENOME
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                                                                                                                                                                                                                                          2170;
                                                                                                                                                                                                                                                                                                                                                                                                                                                SCI
                                                                                                                                                                                                                                                                                                                                                                                                      Barash
                                                                                                                                                                                                                                          262pp;
                                                                                                                                                                                                                                                                                                                                                                                                      SC
                                                                                                                                                                                                                                          English
                  disease), cognitive disorders
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gkin

disorders

(e.g., psoriasis),

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RESULT 1:
ADJUNT/RESULT 1:
ADJUNT/RESULT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      맑
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Best Local S
Matches 45
31-JAN-2000

04-FEB-2000

24-FEB-2000

16-MAR-2000

17-MAR-2000

18-AFR-2000

19-MAY-2000

07-JUN-2000

07-JUN-2000

07-JUL-2000

07-JUL-2000

11-JUL-2000

14-AUG-2000

14-AUG-2000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            breast neoplasms; liver neoplasm; cardiovascular disorder; cardiac arrest; cerebrovascular disorder; cerebral ischaemia; angiogenesis; nervous system disorder, Alzheimer's disease; i ocular disorder; corneal infection; wound healing; epithelial cell proliferation; skin aging; sunburn;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                26-AUG-2002;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     08-JAN-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            organ transplantation; cell culture; tissue regeneration; chemotaxis; food additive; preservative; cardiovascular system associated antigen; nuclear factor kappaB; NFkappaB; promoter element; human; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     autoimmune
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human cardiovascular system associated polypeptide-related DNA SeqID2170.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ADJ08782
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Sequence 17581 BP; 4762 A; 3663 C; 4018
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          2898
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        4841 GGATCACCTGAGGCCAGGAGTTCGAGACCCAGCCTGGCCAACATAG
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               45;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     disease;
2000US-0179065P.
2000US-0184664P.
2000US-0184664P.
2000US-018465P.
2000US-01990769.
2000US-01990769.
2000US-0295515P.
2000US-0214886P.
2000US-0214886P.
2000US-0215135P.
2000US-0217487P.
2000US-0217487P.
2000US-0217487P.
2000US-0217487P.
2000US-0218290P.
2000US-0218290P.
2000US-022513P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225266P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      DNA; 17581
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            rheumatoid arthritis; hyperproliferative disorder;
iver neoplasm; cardiovascular disorder;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Score 45; DB Pred. No. 2.9
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2.9e-10
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          infection;
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The invention relates to an isolated nucleic acid molecule encoding a CC human cardiovascular system associated polypeptide (or antigens), or its C fragment. Also included recombinant vectors, recombinant host cells, an CC isolated human cardiovascular system associated polypeptide (including CC its fragment, allelic variant, species homologue or epitope), an isolated CC antibody that binds specifically to a human cardiovascular system CC susceptibility to a pathological condition or CC susceptibility to a pathological condition comprising determining the CC presence or absence of a mutation in human cardiovascular system CC associated nucleic acid and diagnosing a condition based on the presence or absence of the mutation), identifying a binding partner to human CC cardiovascular system associated polypeptides, the gene corresponding to the human cardiovascular system associated polypeptides, the gene corresponding CC an activity in a biological assay comprising expressing the human CC cardiovascular system associated cDNA in a cell, isolating the cardiovascular system associated cDNA in a cell, isolating the corresponding to condition the supernatant, detecting an activity in a biological assay and identifying cardiovascular system associated cDNA in a cell, isolating the cardiovascular system associated nucleic acids and polypeptides are used the prevent, treat or ameliorate a medical condition (for example in
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14-AUG-2000

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep), example autoimmune diseases such as rheumatoid arthritis, hyperproliterative disorders, for example neoplasms of the breast or liver, cardiovascular disorders, for example cardiac arrest,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       wound healing; skin aging; organ transplantation; tissue regeneration; anti-infertility.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                               17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Cardiovascular system antigen; human; mouse; rabbit; goat; horse;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Human cardiovascular system antigen genomic DNA SEQ ID No 2312.
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2000US-0184664P.
2000US-0184659P.
2000US-019076P.
2000US-0199076P.
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S-0246532P.
S-0246609P.
S-0246610P.
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Sequences AAS35741-AAS36942 represent genomic DNA molecules, which encode CC the cardiovascular system antigen polypeptides of the invention. CC Cardiovascular system antigen and their associated polynucleotides are useful in the diagnosis, treatment and prevention of various types of CC disorders in e.g. humans, mice, rabbite, goats, horses, cats, dogs, CC chickens or she pathological condition can be determined by CC detecting the presence or absence of a mutation in a cardiovascular System antigen polynucleotide. The treatable disorders include autoimmune CC diseases such as rheumatoid arthritis, hyperproliferative disorders such as neoplasms of the breast or liver, cardiovascular disorders such as cardiac arrest, cerebrovascular disorders such as cerebral ischaemia, CC cardiac arrest, cerebrovascular disorders such as cerebral ischaemia, CC cardiac arrest, viruses and fungi, ocular disorders such as corneal correctoria, viruses and fungi, ocular disorders such as corneal correctoria, endocrine disorders such as premature labour and infertility, C gastrointestinal disorders such as premature labour and infertility, C gastrointestinal disorders such as Crohn's disease, renal disorders such as glomerulonephritis and respiratory disorders such as asthma and CC gastrointestinal disorders such as Crohn's disease, renal disorders such cas glomerulonephritis and respiratory disorders such seath and the printed sequence data for this patent did not form part of the printed CC sequence data for this patent did not form aart of the printed CC sequence data for this patent did not form aart of the printed CC at fitp.wipo.int/pub/published_pct_sequences
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                                                                                                                                                                                                                                                                                                                                                                                                                                                          cardiovascular
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Query Match 1.4%; Score 45; DB 4; Length 17946; Best Local Similarity 100.0%; Pred. No. 2.9e-10;
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RESULT 151

ABA15608

ID ABA156

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; vulnerary;
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RESULT 152
ADE47506
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                                                                                                                                                                                                                                                                                                                                                                               The invention relates to novel genes (ABA11004-ABA21534) and proteins CC (ABB14678-ABB18001) useful for preventing, treating or ameliorating comedical conditions e.g. by protein or gene therapy. The genes are CC isolated from a range of human tissues disclosed in the specification. CC The nucleic acids, proteins, antibodies and (ant) agonists are useful in CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast and CC ovarian cancer and other cancers of the adrenal gland, bone, bone marrow, CC breast, gastrointestinal tract, liver, lung, or urogenital; (b) immune CC disorders e.g. Addison's disease, allergies autoimmune haemolytic CC anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease, CC multiple sclerosis, rheumatoid arthritis and ulcerative colitis; (c) CC cardiovascular disorders such as myocardial ischaemias; (d) wound healing CC infections. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly CC from WIPO at ftp.wipo.int/pub/published_pct_sequences
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Matches 45
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                                                                                                                                                                                                                                                                                                                                             Sequence 17946 BP; 4718 A; 3994 C;
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                       29-JAN-2004
                                                             ADE47506;
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                                                                                                                                                                                              GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG
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J. 2.9e-10;
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2000US-022950PP
2000US-023043PP
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2000US-0231268P
2000US-0231968P
2000US-0232399P
2000US-0232399P
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human cardiovascular system related genomic Daw, HAUVA. human, cardiovascular system related physperidic cancer; proliferative disorder, footal abnormality, developmental abnormality, haematopositic disorder, TADE; autohuma disease; rheumatod arthritis; miflammation; allergy; neurological disorder; Alzheimer's disease; parkinson's disease; cognitive disorder; Ralzheimer's disease; skin disorder; purological disorder; Ralzheimer's disease; endocrine disorder; purological disorder; Alzheimer's disease; endocrine disorder; purological disorder; Alzheimer's disease; endocrine disorder; purological disorder; Alzheimer's endocrine disorder; purological disorder; endocrine disorder; purological disorder; postrolitestinal disorder; penglar, penglar, disorder; penglar, penglar, disorder; penglar, penglar, penglar, disorder; penglar, penglar, dis	
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RESULT 153
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DE GAGLAR
KW Cardak
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08-DEC-2000;

08-DEC-2000;

08-DEC-2000;

08-DEC-2000;

11-DEC-2000;

11-DEC-2000;
                                                                       autoimmune disease; rheumatoid arthritis; hyperproliferative disorder; breast neoplasms; liver neoplasm; cardiovascular disorder; cardiac arrest; cerebrovascular disorder; cerebral ischaemia;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             New cardiovascular system related polynucleotides and polypeptides, useful for preventing, treating, or ameliorating a medical condition, such as cancer of cardiovascular tissues and cancer metastases.
               angiogenesis; nervous system disorder; Alzheimer's disease; ocular disorder; corneal infection; wound healing; epithelial cell proliferation; skin aging; sunburn;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence 17946 BP; 4718 A; 3994 C;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      genomic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Claim 1; SEQ ID NO 2312; 262pp; English.
                                                                                                                                                    Human cardiovascular system associated polypeptide-related DNA SeqID2312
                                                                                                                                                                                           04-NOV-2004
                                                                                                                                                                                                                               ADJ08924;
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 transplantation;
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2000US-0251719P.
2000US-0251479P.
2000US-0251868P.
2000US-0251868P.
2000US-0251869P.
2000US-0251999P.
2000US-0254990P.
2000US-0254990P.
2001US-0259678P.
2001US-00764869.
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Pred. No.
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tissue regeneration; chemotaxis;
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2.9e-10;
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                                                         infection;
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18-APR-2000;
19-MAY-2000;
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                                                       2000US-0232080P.
2000US-0232081P.
2000US-0231968P.
2000US-0232397P.
2000US-0232399P.
2000US-0232399P.
2000US-0232400P.
2000US-0232400P.
2000US-0232401P.
2000US-0233063P.
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2000US-0198123P.
2000US-0205515P.
2000US-0214886P.
2000US-0214886P.
2000US-0214886P.
2000US-0214880P.
2000US-0215880P.
2000US-0218290P.
2000US-0224518P.
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2000US-0225213P.
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2000US-023544P.
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RESULT 154
ACM45138
ID ACM451
XX ACM451
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XX Human
DX Human
XX Cytost
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         The invention relates to an isolated nucleic acid molecule encoding a CC fumman cardiovascular system associated polypeptide (or antigens), or its CC fragment. Also included recombinant vectors, recombinant host cells, an CC isolated human cardiovascular system associated polypeptide (including its fragment, allelic variant, species homologue or epitope), an isolated CC antibody that binds specifically to a human cardiovascular system CC associated polypeptide, diagnosing a pathological condition or CC susceptibility to a pathological condition or CC susceptibility to a pathological condition based on the presence CC presence or absence of a mutation), identifying a binding partner to human CC cardiovascular system associated polypeptides, the gene corresponding to CC che human cardiovascular system associated cDNA sequence and identifying CC an activity in a biological assay comprising expressing the human CC cardiovascular system associated cDNA in a cell, isolating the corresponding to CC the protein in the supernatant having the activity. The human CC cardiovascular system associated mucleic acids and polypeptides are used to prevent, treat or ameliorate a medical condition (for example in CC humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep), for cxample autoinmune diseases such as theumacoid arthritis, chyperproliferative disorders, for example neoplasms of the breast or liver, cardiovascular disorders, for example cardiac arrest.
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                                                                                                                                                                                                                                                                                                                                                                  ACN45138;
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                                                                                          28-FEB-2003; 2003WO-US006235.
                                                                                                                                  12-SEP-2003
                                                                                                                                                                      WO2003073826-A2
                                                                                                                                                                                                           Homo sapiens.
                                                                                                                                                                                                                                               Cytostatic; carcinoma; lymphoma; cancer; human;
                                                                                                                                                                                                                                                                                    Human genomic sequence hCG17175.
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                   (SAGR-) SAGRES DISCOVERY
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Disclosure; SEQ ID NO 2312; 262pp; English.
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2002US-00091504.
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The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bloactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bioactive agent capable of modulating the activity of CAP; (iv) for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Claim 1; SEQ ID NO 1936; Opp; English
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                                                                                                                                                                                                                                                                                                         Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
                                                                                                                                                                                                                                                                                                                                                                              WPI; 2003-328604/31.
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                                                                                                                                                                                                        Claim 1; SEQ
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Pred. No.
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                       CC associated (CA) nucleic acid, comprising two or more nucleic acid probes.

CC rhe invention also relates to a peptide array comprising two or more solution also relates to a peptide array comprising two or more composed by a CA nucleic acid sequence, a compound composed that binds to a polypeptide, which is prepared by immunizing a host animal composition comprising the polypeptide or its fragment which composition comprising the polypeptide or its antigen binding composition comprising the polypeptide or its antigen binding composition comprising the antibody and a carrier, a method of fragment, a composition comprising the composition of actions, a composition comprising the composition of detecting a CA nucleic acid, a method of diagnosing cancer, a composition of a CA nucleic acid, a method of inhibiting expression of a CA nucleic acid in a cell. The CA nucleic acids are useful for detecting CA composition acids. The antibody is useful for detecting the presence or cancer cells in an individual which involves contacting cells from the individual with the antibody and detecting a complex of a CA protein from the cancer cells and the antibody, where the detection of the complex of the complex center when the cancer cells and the antibody, where the detection of the complex center when the cancer cells and the antibody, where the detection of the complex center when the cancer cells and the antibody, where the detection of the complex center when the cancer cells and the antibody.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 775; 198pp; English
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Nucleic acid array useful for detecting cancer associated nucleic acid,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              WPI; 2005-273395/28
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               The invention relates to a nucleic acid array for detecting a cancer
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45; Conserv
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           correlates with the
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Pred. No.
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individual. The composition is useful for inhibiting growth of cancer

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RESULT 157
ACN45014
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Best Local S
Matches 45
                               Query Match
Best Local Similarity
Matches 45; Conserv
                                                                                                                                        The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bioactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bioactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing carcinoma; (vii) for inhibiting the activity of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for neutralizing the effect of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the CA genee are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent respectation.
                                                                                                                                                                                                                                                                                                                                                                                                                Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human genomic sequence hCG14907
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ACN45014 standard;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     cells in an individual or for delivering a therapeutic agent to cancells in an individual. The invention is also useful for diagnosing cancer, for treating cancer and for inhibiting expression of a CA ge a cell. This sequence represents human cancer-associated genomic DNF
                                                                                               Sequence 32706
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   01-MAR-2002; 2002US-00087192
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    28-FEB-2003; 2003WO-US006235
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     12-SEP-2003
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Homo sapiens
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                                                                                                                               US2002182586A1,
                                                                                                                                                                                                                                                                                                                                                                                Claim 1; SEQ ID NO 1750; Opp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    (SAGR-)
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2898 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 2942
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    2003-328604/31.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    SAGRES
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       GTGCCACTGCAGTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 7181
                                Conservative
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                                                                                               BP; 8225 A;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    DISCOVERY
                                                                                                                              for which no sequence data was published
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     DNA;
                                            1.4%;
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100.0%; Pred. No. 2.8e-10;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       32706
                                                                                               7861 C; 8277 G; 8343 T; 0 U; 0 Other;
                               0,
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                                              Score 45;
Pred. No.
                                Mismatches
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                                              DB 11; I
2.8e-10;
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                               0,
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                                                            Length 32706;
                                Indels
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                               0,
                               Gaps
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RESULT 158
ADL82795/c
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Matches 45; Conservation
                                                                                                                                                                                                                                                                                                                                                        The invention relates to a method of inhibiting the proliferation of a cancer cell comprises contacting the cell with a semaphorin3B (SEMA3B) polypeptide. The composition and methods are useful in diagnosing or treating cancer. The SEMA3B polypeptide inhibits tumour growth and induces apoptosis in cancer cells. The present sequence represents DNA encoding human semaphorin3B, SEMA3B.
                      SNP detection; diagnosis; non-insulin dependent diabetes; antidiabetic; anorectic; endocrine disease; gastrointestimetabolic disorder; nutritional disorder; gene; ds.
                                                                                                     08-SEP-2005
                                                                                                                                                         AEB32373
                                                                                                                                                                                                                                                                                                                                  Sequence 36534 BP; 7493 A; 10597 C; 10438 G; 8006 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                       Disclosure; SEQ ID NO 3; 75pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Inhibiting the proliferation of a cancer cell (e.g. breast cancer cell, lung cancer cell or prostate cancer cell) comprises contacting the cell with a semaphorin3B polypeptide that suppresses tumor growth.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 31-OCT-2001; 2001US-0335783P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            31-OCT-2002; 2002US-00285351.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     apoptosis; human;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   cancer cell proliferation; semaphorin3B; SEMA3B; cancer; tumour growth;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human semaphorin3B,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        20-MAY-2004
                                                                         Human genomic DNA #14.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Minna J,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 US2003166557-A1
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                                                                                                                                                                                                                           7478
                                                                                                                                                                                                                                                    3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        2003-898098/82.
DB; ADL82793.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG
                                                                                                                                                         standard; DNA; 38678
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Tomizawa Y,
                                                                                                     (first entry
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (first
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                                                                                                                                                                                                                                                                                           Score 45;
Pred. No.
                                                                                                                                                         BP.
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                                                                                                                                                                                                                                                                                                      DB 11;
                                                                                                                                                                                                                                                                                           2.8e-10;
                                     gastrointestinal
                                                                                                                                                                                                                                                                                                       Length 36534;
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                                       disease;
                                                                                                                                                                                                                                                                              Gaps
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Homo sapiens

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acid molecule for detecting at least one SNP given in the specification, an isolated nucleic comprising at least about 12 contiguous nucleotides, genotyping at least cone SNP position given in the specification in a sample, identifying an individual having or at risk of developing a disorder and a kit comprising at least one containing the detection reagent.

CC Determining whether a trait is linked to one of the human chromosomes or its sub-region comprises determining whether the trait is linked to one or more SNPs using the detection reagent. Senotyping at least one SNP position given in the specification in a sample for allele stated to suppose that differentiates between alternative determining which allele is present at the at least one SNP position. CC determining which allele is present at the at least one SNP position. CC nucleic acid sample from the individual. The disorder is type II diabetes or obesity. The detection reagent is consider, particularly type II diabetes or obesity. This sequence creptesents human genomic DNA used in the scope of the invention. Note: The sequence data for this patent did not form part of the printed consideration but was obtained in electronic format from USPTO at the constraint of the printed constraints.
ARBSULT 160
ARB32391/c
ID ARB323
XX
AC ARB323
XX
DT 08-SEP
XX
DE Human
XX
XX
DX SNP de
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                                                                                                                                                                                                                                                                                                               Matches
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Best Local 9
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The invention relates to a detection reagent capable of detecting one more single nucleic acid polymorphisms. The invention also relates to determining whether a trait is linked to one of the human chromosomes
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   New detection reagent capable of detecting 1, 100, 500, 1000 or 5000 or more single nucleic acid polymorphisms, useful in identifying an
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 08-SEP-2000; 2000US-0231397P.
10-SEP-2001; 2001US-00948947.
                                   Human genomic DNA #32.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Disclosure; SEQ ID NO 136; 31pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            US2005147987-A1.
SNP detection; diagnosis; non-insulin dependent diabetes; obesity;
                                                                        08-SEP-2005
                                                                                                                                                ABB32391 standard; DNA; 38684
                                                                                                                                                                                                                                                                                                                                                                              Sequence 38678 BP; 9340 A; 9040 C; 9074 G; 10537 T; 0 U; 687 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    its sub-region, a computer readable medium having stored in it relational information given in the specification, an isolated
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                individual having or at risk of developing type II diabetes or obesity.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               (APPL-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     19-JUL-2004; 2004US-00893315
                                                                                                                                                                                                                                        23579
                                                                                                                                                                                                                                                                         2895 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 2939
                                                                                                                                                                                                                                                                                                               45
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                                                                                                                                                                                                                                                                                                                              Similarity
                                                                                                                                                                                                                                        GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 23535
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Zhang JN,
                                                                                                                                                                                                                                                                                                               Conservative
                                                                       (first entry)
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                                                                                                                                                                                                                                                                                                                            100.0%;
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                                                                                                                                                                                                                                                                                                           0
                                                                                                                                                                                                                                                                                                                            Score 45;
Pred. No.
                                                                                                                                                                                                                                                                                                               Mismatches
                                                                                                                                                                                                                                                                                                                                                DB 14;
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                                                                                                                                                                                                                                                                                                                                             Length 38678;
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                                                                                                                                                                                                                                                                                                           Gaps
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ADN31618
ID ADN3
XX
AC ADN3
XX
DT 12-1

ADN31618 standard; DNA; 39566 BP

12-AUG-2004 ADN31618;

(first entry)

RESULT 161

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CC relational information given in the specification, an isolated nucleic comprising at least about 12 contiguous nucleotides, genotyping at least one SNP position given in the specification in a sample, identifying an comprising at least one container containing the detection reagent. CC comprising at least one container containing the detection reagent. CC comprising at least one container containing the detection reagent. CC its sub-region comprises determining whether the trait is linked to one or more SNPs using the detection reagents. Genotyping at least one containing whether the trait is linked to one co more SNPs using the detection reagents. Genotyping at least one SNP co position given in the specification in a sample comprises contacting the calleles at at least one SNP position given in the specification, and cC determining which allele is present at the at least one SNP position. CC comprises genotyping at least one SNP given in the specification, and cC determining which allele is present at the at least one SNP position. CC comprises genotyping at least one SNP given in the specification in a sample comprise disorder is type II diabetes (non-insulin dependent diabetes) or obesity. The detection reagent is useful in identifying an individual having or at risk of developing a disorder particularly type II diabetes or obesity. This sequence cC represents human genomic DNA used in the scope of the printed consideration but was obtained in electronic format from USPTO at
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                                                                                                                            Query Match
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          New detection reagent capable of detecting 1, 100, 500, 1000 or 5000 or more single nucleic acid polymorphisms, useful in identifying an individual having or at risk of developing type II diabetes or obesity.
                                                                                                                                                                       Sequence 38684 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     more single nucleic acid polymorphisms. The invention also relates to determining whether a trait is linked to one of the human chromosomes its sub-region, a computer readable medium having stored in it the SNP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    The invention relates to a detection reagent capable of detecting one or more single nucleic acid polymorphisms. The invention also relates to
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 154; 31pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Venter JC, Zhang JN,
Naik A, Subramanian G,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                08-SEP-2000; 2000US-0231397P.
10-SEP-2001; 2001US-00948947.
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metabolic disorder; nutritional disorder; gene; ds.
                                                                                                                                                                                                                  seqdata.uspto.gov/sequence.html.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                19-JUL-2004; 2004US-00893315
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                                                                                     Local Similarity nes 45; Conserv
23584 GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 23540
                                          2895
                                        GGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACA 2939
                                                                                     Conservative
                                                                                                                                                                         9340 A; 9042 C;
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lage T;
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. 2.8e-10;
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RESULT 162
ABX14652
ID ABX146
XX ABX146
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Best Local :
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                                                                                                                                                                                                                                                                                                                                                 The invention relates to a new compound 8-80 nucleobases in length (an antisense oligonucleotide) targeted to a nucleic acid molecule encoding squalene synthase (also known as farnesyl diphosphate farnesyl transferase 1), where the compound specifically hybridises with the nucleic acid molecule encoding human squalene synthase appearing as ADN31611 and inhibits the expression of squalene synthase in cells or tissues, screening for a modulator of squalene synthase, a diagnostic method for identifying a disease state, a kit or assay device comprising the compound and treating an animal having a disease or condition associated with squalene synthase. The compound and methods are useful in diagnosing and treating disorders related to cholesterol biosynthesis e.g. atheroselerosis, coronary heart disease and hypercholesterolaemia. The present sequence is a squalene synthase genomic DNA sequence, a target for the artisense alignmentations.
                                                                  Human; ds; gene; squalene synthase; cholesterol-related disease; cardiovascular disease; chromosome 8.
                                                                                                   Human gene encoding squalene synthase.
                                                                                                                         05-MAR-2003
                                                                                                                                                                                                                                                                                                                     Sequence
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                New oligonucleotide targeted to a nucleic acid molecule encoding squalene synthase, useful in diagnosing and treating atherosclerosis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             23-NOV-2002; 2002US-00304125
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human; ds; antisense; farnesyl diphosphate i atherosclerosis; coron
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Human squalene
                                           Homo sapiens
                                                                                                                                                ABX14652;
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            variation
                                                                                                                                                                      ABX14652 standard;
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                                                                                                                                                                                                                                                                                    Local
                                                                                                                                                                                                                              25338
                                                                                                                                                                                                                                                                                                                                             the antisense oligonucleotides.
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                                                                                                                                                                                                                                                                                                                      39566 BP;
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Location/Qualifiers replace(825,A) /*tag= o
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hate farnesyl transferase 1; cholesterol;
coronary heart disease; hypercholesterolaemia.
                                                                                                                                                                      DNA; 40090
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Pred. No.
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9054. .25147
/*tag= g
/number= 3
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replace(8310,A)
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replace(7212,G)
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/*tag= c
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replace(8462,G)
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replace(7398,C)
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replace(7355,T)
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replace(4887,T)
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replace(4791,T)
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2058. .37739
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                                                       eplace(8873,T)
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replace (29761,T)
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replace(10460,T)
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replace(9310. .9312,GA)
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/*tag= k
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/standard_name=
replace(23963,T)
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replace(20362,A)
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replace(26018,G)
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replace(22230,T)
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eplace(26625,G)
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eplace(26078,A)
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FH CDS
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Best Local Similarity
Matches 45; Conserv
                                                                                                         disease diagnosis; gene eapression associated enzyme peptide; human; gene; ds.
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/standard_name= "
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replace(34532,C)
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replace(34179,T)
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2058. .37739

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2058. .2156
/*tag= b
/number= 1
                                                                                                replace (825, A)
                                                                        "Single nucleotide
                                                                         polymorphism"
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DNA.

disorder;

gene

expression;

FT / ray- and / ray- ray- ray- ray- ray- ray- ray- ray-	variation	variation /	/*tag= /standar intron 90542	variation r	/*tag= /standar exon 8870s	variation /	/*tag= t /standard_na riation replace(8310	/*tag: /numbe variation replace	FT /*tag= r /*single nucleotide polymorphism" FT intron 80958869	/*tag= 2 /number= 2	FT /*tag= p FT /standard name= "Single nucleotide polymorphism" T7007 8004	variation	variation replace	e (variation replace	variation /	variation x	variation r	variation r	variation r	variation r	variation	FT intron 21577996 FT /*tag= d FT /number= 1 FT variation replace(2632,T)
variation /	intron 3079337517 /*tag= ay /number- 6	variation replace (30732,C) /*tag= ax /*tag= name- "	FT exon 3064030792 FT exon 4.tag= aw FT /*tag= aw FT /*tag= aw	/#Caradard_name= "Single variation replace(29761,T) /*tag= av /*ta	variation replace(29572,T) /*tag= au	intron	/standar 29366. /*tag=	/standar variation replace /*tag=	FI /"tag= ap FT variation replace(28032,A) FT /*tag= ag	variation replace	/standa variation replace /*rar=	variation replace(26078,A) /*tag= an	variation replace(26018,G) /*tag= am /*tag= am	variation replace (25686, A) /*tag= al /atandard name-	intron 253402	/standard 251482 exon /*tag= a	/standard name= "Single variation replace(23963,T) /*tag= ai	/standa: variation replace /*tag=	/standa: variation replace /*tag=	variation r	/standard_name= "Single variation replace(21166,A) /*tag= ae	/standard_name= variation replace(20362,A) /*tag= ad	FT /*tag= ab FT /standard name= "Single nucleotide polymorphism" FT variation replace(20204,G) FT /*tag= ac

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RESULT 164
ADZ13149/c
ID ADZ131
XX ADZ131
XX ADZ131
XX DIAGNO
XX DIAGNO
XX DIAGNO
XX DIAGNO
XX DIAGNO
XX DOZOOS
XX HOMO 8
XX WO2005
XX WO2005
XX JONAPR
XX 23-SEP
XX CHIR
XX ICHIR
XX WPI; 2
XX WPI; 2
XX WPI; 2
XX WNuclei
PT compri
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Best Local Similarity
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            Nucleic acid array useful for detecting cancer associated nucleic acid, comprises two or more nucleic acid probes.
                                                 WPI; 2005-273395/28.
                                                                          Morris
                                                                                                                        23-SEP-2003; 2003US-00669920
                                                                                                                                               23-SEP-2004; 2004WO-US031617.
                                                                                                                                                                        07-APR-2005.
                                                                                                                                                                                                 WO2005031001-A2
                                                                                                                                                                                                                       Homo sapiens
                                                                                                                                                                                                                                              cytostatic; gene;
                                                                                                                                                                                                                                                           Diagnosis; DNA microarray; microarray; biochip; cancer; neoplasm;
                                                                                                                                                                                                                                                                                  Human cancer-associated genomic DNA #56
                                                                                                                                                                                                                                                                                                           16-JUN-2005
                                                                                                                                                                                                                                                                                                                                                          ADZ13149 standard; DNA; 57105
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             variation
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                                                                                                 (CHIR ) CHIRON CORP
                                                                                                                                                                                                                                                                                                                                                                                                                                              2898 GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 2942
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                                                                                                                                                                                                                                                                                                                                                                                                                      GGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAG 25757
                                                                          Malandro
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                                                                                                                                                                                                                                                                                                          (first entry
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replace(34179,T)
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replace(32525,G)
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/*tag= bi
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be
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                     0; Mismatches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           nucleotide polymorphism'
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                nucleotide
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Length 40090;
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##666666666666666666666666666666668
                                                                CC The invention also relates to a peptide array comprising two or more consolated polypeptides encoded by a CA nucleic acid sequence, a compound that binds to a polypeptide, which is prepared by immunizing a host animal CC with a composition comprising the polypeptide or its fragment which the acomposition comprising the polypeptide or its antigen binding CC fragment and collecting cells from the host expressing antibodies against the antibody and a carrier, a method of fragment, a composition comprising the CC antibody and a carrier, a method of ecreening for anticancer activity, a complete acid in a cell. The CA nucleic acid, a method of disgnosing cancer, a cC nucleic acid in a cell. The CA nucleic acids are useful for detecting CA concleic acids. The antibody is useful for detecting the presence or cancer cells in an individual which involves contacting cells from the individual with the antibody, where the detection of the complex correlates with the presence of cancer cells in the composition is useful for inhibiting growth of cancer cells in an individual. The invention is also useful for day agent to cancer cells in an individual. The invention is also useful for day agent to cancer the cell. This sequence represents human cancer-associated genomic DNA of the coll. This sequence represents human cancer-associated genomic DNA of
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        The invention relates to a nucleic acid array for detecting associated (CA) nucleic acid, comprising two or more nucleic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Disclosure; SEQ ID NO 669; 198pp; English
                                                       invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     nucleic acid,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        20
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              acid
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밁 8 39001 3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122

Matches Query Match

Local Similarity nes 45; Conserv

Conservative 0;

1.4%;

Score 45; DB 14; pred. No. 2.8e-10; Mismatches

Length 57105; Indels

0;

0

Gaps

0

Sequence 57105 BP; 15389 A; 12942 C; 12984 G; 15770 T; 0 U; 20 Other;

RESULT 165
ABK83563
ID 83563
XX ABK835
XX ABK835
XX Human
XX Human;
KW Human;
KW Viral
KW rheuma
KW rheuma
KW Grohn;
KW Grohn; Human cDNA differentially expressed in granulocytic cells #134. ABK83563 ABK83563 14-AUG-2002 standard; cDNA; 57248 (first entry)

Human; 88; granulocytic cell; DNA chip; bacterial infection; viral infection; parasitic infection, protozoal infection; fungal infection; sterile inflammatory disease; psoriasis; rheumatoid arthritis; glomerulonephritis; asthma; thrombosis; cardiac reperfusion injury; renal reperfusion injury; ARDS; adult respiratory distress syndrome; inflammatory bowel disease; Crohn's disease; ulcerative collitis; periodontal disease; granulocyte activation; chronic inflammation; allergy.

Homo sapiens.

WO200228999-A2.

11-APR-2002.

03-OCT-2001; 2001WO-US030821.

03-OCT-2000; 2000US-0237189P

GENE LOGIC INC.

Beazer-Barclay Y, Weissman Z, Yamaga ຸດ

WPI; 2002-435328/46

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CC DNA chip analysis as given in the specification, and comparing the expression level to an expression of GCA. Also included are CC differential expression of Gs is indicative of GCA. Also included are CC expression of at least one gene in Gs; (2) screening (M3) for an agent cc expression of at least one gene in Gs; (2) screening (M3) for an agent CC expression of at least one gene in Gs; (2) screening (M3) for an agent cc tissue, an allergic response in a subject, exposure of a subject to a CC pathogen or sterile inflammatory disease using the gene expression CC pathogen or sterile inflammatory disease using the gene expression cc pathogen or sterile inflammatory disease, by detecting the level of expression in a sample of the tissue of gene(s) from Gs, where the level of expression of the gene is indicative of inflammation; (4) treating CC (M5) an inflammation (especially chronic) or in a tissue, an allergic cresponse in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by contacting a tissue having inflammation with an CC agent that modulates the expression of gene(s) from Gs in the tissue. M1 is useful for detecting an agent capable of modulating GCA preferably in an CC (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammation with an CC (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammation with an CC (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammation with an CC (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile GCA preferably in an Inflammation of the preferably in an Inflammation of the M10 in the tissue, and the control of the M10 in the tissue, and the control of the promoted at the control of the state of the promoted at the control of the state of the control of the state of the control 
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Best Local S
17-FEB-2004; 2004WO-US004730.
                                                                          02-SEP-2004
                                                                                                                                                    WO2004074320-A2
                                                                                                                                                                                                                                                                                        Human; ds; cancer-associated protein; gene; cytostatic; cancer;
leukaemia; lymphoma; CAP.
                                                                                                                                                                                                                                                                                                                                                                                                              Human cancer-associated genomic DNA HD18-038
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        18-NOV-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ABD32902
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        ABD32902 standard; DNA; 65277
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 57248 BP; 15003 A; 13601 C; 13307
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Claim 1; SEQ ID NO 134; 114pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Detecting granulocyte activation by detecting differential expression of genes associated with granulocyte activation, which serves as diagnostic markers that is useful for monitoring disease states and drug toxicity.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   2888 TGAGGCAGGTGGATCACCTGAGGCCAGGAGTTCGAGACCAGCCTG 2932
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                45;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Length 57248;
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RESULT 167 ACN43986/c ID ACN439

ACN43986 standard; DNA; 73995 BP

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GATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAGC 27364 GATCACCTGAGGCCAGGAGTTCGAGACCAGCCTGGCCAACATAGC 2943 Matches Query Match

Local Similarity

1.4%;

Score 45; Pred. No. Mismatches

DB 13; 2.8e-10;

Length 65277; Indels

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Gaps

0

Conservative

0,

Sequence 65277 BP;

19651 A; 11706 C; 12664 G; 21256 T; 0 U; 0 Other;

SARARA

18-NOV-2004

(first entry)

Human genomic sequence hCG40211.

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CC mentioned nucleotide sequences, an isolated polypeptide (encoded within CC an open reading frame of a CA sequence selected from any of the 95 CC polynucleotide sequences as mentioned in the specification, or its CC polynucleotide sequences as mentioned in the specification, or its CC complement), an isolated antibody, (or its antigen binding fragment) that binds to the above polypeptide, a hybridoma that produces the above CC monoclonal antibody, a pharmaceutical composition comprising the above CC antibody and a pharmaceutical excipient, a kit for detecting cancer CC cristody and a pharmaceutical excipient, a kit for detecting cancer CC or for detecting the antibody cited above, methods for diagnosing cancer CC individual, a method for inhibiting growth of cancer cells in an CC individual, a method for delivering a therapeutic agent to cancer cells in an individual, a method for thibiting growth of cancer cells in an individual, an electronic library comprising the above CC polynucleotide or polypeptide (or their fragments), methods of screening CC individual, a method for for a bioactive agent capable of modulating CC individual, a method for for a bioactive agent capable of modulating CC individual, a method for for a bioactive agent capable of modulating CC individual, and the composition and methods for detecting cancer CC associated with expression of a polypeptide in a test cell sample, a composition and methods are useful for detecting, diameter in a cell. The composition and methods are useful for detecting,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          14-FEB-2003;
14-MAR-2003;
15-APR-2003;
13-JUN-2003;
15-SEP-2003;
                                                                     diagnosing, preventing and treating cancers, especially lymphoma and leukaemia. These may also be used in screening for agents that modula cancer. The present sequence is a human CAP genomic sequence. Note: I sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WI specification, but was obtained in electronic format directly from WI
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             in the specification, or its complement. The nucleic acids encode cancer associated proteins. Also included are an expression vector comprising the isolated nucleic acid cited above, a host cell comprising the above recombinant nucleic acid or expression vector, a microarray for detectif a cancer-associated (CA) nucleic acid comprising at least one probe comprising at least 10 contiguous nucleotides of any of the above-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      The invention relates to an isolated nucleic acid comprising at least 10 contiguous nucleotides of any of the 233 polynucleotide sequences given in the specification, or its complement. The nucleic acids encode cancer
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    claim 16; seqid 602; 310pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      New isolated cancer-associated polynucleotides and polypeptides useful for diagnosing, preventing or treating cancers, especially lymphoma an leukemia, or in screening for agents that modulate cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         (SAGR-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   15-DEC-2003;
                                              ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 DW,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       ; 2003US-00367094.
; 2003US-00388838.
; 2003US-00417375.
; 2003US-00461862.
; 2003US-00663431.
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                                                                                 from WIPO
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RESULT 168
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Best Local (
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   are associated with carcinomas. The sequences are useful for: (1) for screening drug candidates; (ii) for screening of bloactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of bloactive agent capable of modulating the activity of CAP; (iv) for a bloactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating carcinoma; (vii) for nuturalizing the effect of CAP; (ix) as a blochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the
                                                                                                                                                                                                                                                                                                    Human; chronic inflammatory joint disease; infection; tumour; antiinflammatory; cytostatic; antiarthritic; antirheumatic; immunosuppressive; gene therapy; etiological pathogenicity; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             uetermining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent US200218258A31 for which the common of the control of th
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           The present invention relates to are associated with carcinomas. ?
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             ACA64942;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Sequence 73995 BP; 17594 A; 18402 C; 19247 G; 18752 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 1; SEQ ID NO 208; Opp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             comprises a
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     28-FEB-2003; 2003WO-US006235
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30-MAY-2001; 2001DE-01027572
                                                                                                                                                                                                                                                     Homo
                                                                                                                                                                                                                                                                                                                                                                                                                             Human FRAP1 DNA corresponding to AL049659.
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                                                            30-MAY-2001; 2001DE-01027572.
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                                                                                                                                                                                                                                                 sapiens.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             entry)
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100.0%; Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 DB 11; : 2.8e-10;
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RESULT 169
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XX AD0794
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KW CPtoSt
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KW CAPC1
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KW 28104(
KW 28104(
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            This invention describes a novel reagent for diagnosis, molecular definition and therapy of chronic inflammatory joint diseases, and other inflammatory disorders, infective or tumour diseases in humans. The products of the invention have antiinflammatory, cytostatic, antirheumatic and immunosuppressive activity and can be used for gene therapy. The reagent of the invention and any proteins and cantibodies derived from it, are used (i) for analysing tissue and blood samples for medical diagnosis; (ii) for diagnosis and characterisation of chronic joint diseases, on the basis of molecular characterisation, and determining the etiological pathogenicity principle of as yet uncharacterised inflammatory diseases, also monitoring progression and/or treatment of disease, and optimisation of therapy and (iii) for developing treatments for inflammatory diseases, particularly of joints, infections and tumours. ACA64801-ACA64965 represent human polynucleotides cused in the method of the invention
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DPF3 region,
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                           /standard name= "Single nucleotide polymorphism"
/note= "This SNP is described as a A/G SNP"
10481
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160
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                                                                                                                                                                             /standard name= "Single nucleotide
/note= "This SNP is described as a
                                                                                                                                                                                                                                                                                                    /standard name= "Single nucleotide polymorphism"
/note= "This SNP is described as a A/C SNP"
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Pred. No.
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T/G SNP"
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v vard_name= "Single nucleotide	variation 36254 /*tag= u /standard name= "Single nucleotide /note= "This SNP is described as a	FT variation 35856 FT /*tag= t FT /standard name= "Single nucleotide polymorphism" FT /note= "This SNP is described as a G/A SNP"		variation 3588 /*tag /*stan /note- /rafietion 3589	variation	Variation 4			variation 2	/*tag= /stanc /note= variation 24767	/*tag= /standa /note= variation 24683	/*tag///////////////////////////////////	cariation /	FT /*tag= h FT /*tag= h FT /standard_name= "Single nucleotide polymorphism" FT /note= "This SNP is described as a G/C SNP" FT variation 18694	/*tag= /stanc/ note=		variation	<pre>/standard name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" variation 10676</pre>
	riation 8	variation 8	variation 7	variation 7	FT // FT // FT variation 61	variation 6	variation 5	variation /	variation 5	variation 5	variation 5	variation 4	variation 4	FT Variation 4.	variation	variation	variation 4	variation 4
*tag= an standard name= "Single nucleotide polymorphism" note= "This SNP is described as a T/A SNP"	<pre>'*tag= am 'standard name= "Single nucleotide polymorphism" note= "This SNP is described as a A/G SNP" 19751</pre>	<pre>/*tag= al standard name= "Single nucleotide polymorphism" note= "This SNP is described as a A/C SNP" 15507</pre>	_ 22	*tag= aj standard name= "Single nucleotide polymorphism" note= "This SNP is described as a T/C SNP" 2720	_ 22	"tag= an standard name= "Single nucleotide polymorphism" note= "This SNP is described as a T/C SNP" 0682	ard name= "Single "This SNP is dea	ard name= "Single "This SNP is des	"Single nucleotide P is described as a	standard name= "Single nuclectide polymorphism" note= "This SNP is described as a T/C SNP" 2079 *tag= ae	"Single nucleotide P is described as a	"Single nucleotide P is described as a	. D	'rtag= z' 'standard name= "Single nucleotide polymorphism" 'note= "This SNP is described as a A/G SNP" 3090	"tag= y 'standard name= "Single nucleotide polymorphism" 'note= "This SNP is described as a A/C SNP" 12799	standard name= "Single nucleotide polymorphism" note= "This SNP is described as a G/C SNP" 2593	ard name= "Single "This SNP is des	note= "This SNP is described as a G/A SNP" 10033 "tag= w

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Best Local Sim
Matches 45;
 The invention relates to cancer-associated proteins (CAP) and the cancer-associated (CA) nucleic acids encoding them. The invention also relates to a method for treating cancers involving administering to a patient an inhibitor of CAP, and a method of screening for anticancer activity in a potential drug involving providing a cell that expresses a CA gene, contacting a tissue sample derived from a cancer cell with an anticancer
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Identifying a subject at risk of of polymorphic variations in the which are associated with breast
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          25-NOV-2002; 2002US-0429136P.
24-JUL-2003; 2003US-0490234P.
                                                                                                                                     Novel human cancer associated protein encoded within open reading f of cancer associated gene, useful as targets for diagnosing cancer.
                                                                                                                                                                                                                                                                                                                                                                                                     Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                  Human; cancer-associated protein; CAP; cancer-associated gene; CA; gene;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Human cancer-associated (CA) gene HD07-103.
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                                                                                                          SEQ ID NO
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                                                                                                         706; 182pp; English.
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Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   breast cancer by detecting the presence DLG1, KIAA0783, DPF3 or CENPC1 regions cancer in a nucleic acid sample from a
from a cancer
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3. 2.7e-10;
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RESULT 171
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                                                        The invention relates to cancer-associated proteins (CAP) and the cancer-
CC associated (CA) nucleic acids encoding them. The invention also relates
CC to a method for treating cancers involving administering to a patient an
CC inhibitor of CAP, and a method of screening for anticancer activity in a
CC contacting a tissue sample derived from a cancer cell with an anticancer
CC contacting a tissue sample derived from a cancer cell with an anticancer
CC drug candidate and monitoring the effect of the anticancer drug candidate
CC on expression of the CA gene. The CAP proteins are useful for detecting
CC cancer associated with expression of a CAP protein in a test cell sample
CC and for screening for a bioactive agent capable of modulating the
CC activity of a CAP protein. The CA nucleic acids are useful for diagnosing
CC cancer, involving determining the expression of a CA nucleic acid in a
CC tissue. This sequence represents a human CA gene of the invention. Note:
CC specification, but was obtained in electronic format directly from WIPO
CC are format directly from WIPO
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Matches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Novel human cancer associated protein encoded within open reading frame of cancer associated gene, useful as targets for diagnosing cancer.
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RESULT 173

ABXO8336 03/c
Continuation (4 of 17)
WP Sequence split into
WP Fragment Name
WP ABXO8336 00
WP ABXO8336 00
WP ABXO8336 03
WP ABXO8336 03
WP ABXO8336 06
WP ABXO8336 06
WP ABXO8336 07
WP ABXO8336 07
WP ABXO8336 09
WP ABXO8336 10
WP ABXO8336 11
RESULT 174

AAD53224_2/c
Continuation (3 of 6) of WP Sequence split into 6
WP Fragment Name
WP AAD53224_0
WP AAD53224_2
WP AAD53224_3
WP AAD53224_3
WP AAD53224_4
WP AAD53224_4
WP AAD53224_5
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RESULT 176
ADM97989 03/c
ADM97989 01/c
Continuation (4 of 17) o
WP Sequence split into 1
WP Sequence split into 1
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WP ADM97989 03
WP ADM97989 05
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WP ADM97989 11
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ADJ25985_03/c

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RESULT 178

AEBS5185,03/c
Continuation (continuation)
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Sequence split into
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AEB85185 02
AEB85185 03
AEB85185 04
AEB85185 06
AEB85185 06
AEB85185 06
AEB85185 10
AEB85185 11
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AEB85185 11
AEB85185 11
AEB85185 11
AEB85185 11
AEB85185 12
AEB85185 11
         26-AUG-2004
                            ADQ17592;
                                               ADQ17592 standard;
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AD050281 00
AD050281 01
AD050281 03
AD050281 03
AD050281 04
AD050281 06
AD050281 06
AD050281 07
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Early detection of soft tissue sarcoma comprises determining expression of a gene in a first soft tissue sample and a normal soft tissue sample and comparing the gene expression, also useful in treating soft tissue
                                                                                                                      cytostatic;
                                                                                                                                       Human; breast specific gene; breast cancer; differential expression;
                                                                                                                                                                      Human breast cancer associated coding sequence SEQ ID NO:
                                                                                                                                                                                                     04-DEC-2002
                                                                                                                                                                                                                                                                   ABT10719
                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 116561 BP; 32234 A; 27837 C; 28253 G;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               The invention relates to a novel method for detecting soft tissue sarcoma which comprises obtaining a first soft tissue sample from an individual
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Aziz N,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 26-NOV-2002; 2002US-0429739P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                26-NOV-2003; 2003WO-US038193
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     specification per se but was submitted in CD format by the inventor.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Example 2;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ds; gene; osteopathic; antiinflammatory; antiarthritic; ge joint space narrowing; osteophyte development; joint pain; osteoarthritis; SNP; single nucleotide polymorphism.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Sequence 122748 BP; 32088 A;
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25-APR-2001; 2001US-0286090P.
23-MAY-2001; 2001US-0292517P.
   WPI; 2003-559141/52
                                                           Jones KA,
                                                                                                                                                                                                                                  19-DEC-2002; 2002WO-US041225
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Osteoarthritis-associated polymorphic nucleotide #441.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Claim 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              (GENE-)
                                                                                                                                                                       20-DEC-2001; 2001US-0342603P
                                                                                                                  (INCY-) INCYTE GENOMICS INC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    37876
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 37920
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Nation M,
                                                       Schafer A;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1.4%;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Score 45;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          31056 C; 30547 G;
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2.7e-1
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      gene therapy;
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RESULT 182
ABK84797/c
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         The invention relates to a method of determining susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain comprising identifying whether the individual has at least one polymorphism in a polymuclectide encoding at least one of the protein listed in the specification. The methods, composition and agent are useful for modulating the susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain that is associated with a disease, preferably osteoarthritis. The cell line and the non-human animal are useful for screening for an agent for diagnosing an individual having susceptibility to joint space narrowing and/or osteophyte development and/or joint pain. This sequence corresponds to the polymucleotide encoding a protein listed in the specification. (Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format directly from WIPO at
                                                                                                                                                                                                                                                                                                                                                                                                     viral infection; parasitic infection; protozoal infection; fungal infection; sterile inflammatory disease; psoriasis; rhumatoid arthritis; glomerulonephritis; asthma; thrombosis; cardiac reperfusion injury; renal reperfusion injury; ARDS; adult respiratory distress syndrome; inflammatory bowel disease;
                                                                                                                                                                                                                                                                                                                                                              adult regitatory distress syndrome; inflammatory power under the disease; ulcerative colitis; periodontal disease; Crohn's disease; ulcerative colitis; periodontal disease; crannilocyte activation; chronic inflammation; allergy.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Human; 88; granulocytic cell; DNA chip; bacterial infection;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Human cDNA differentially expressed in granulocytic cells #1368.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ABK84797
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Disclosure; SEQ ID NO 441;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    a protein.
                                                                                                                                                                                                                     03-OCT-2001; 2001WO-US030821.
                                                                                                                                                                                                                                                          11-APR-2002.
                                                                                                                                                                                                                                                                                               WO200228999-A2
                                                                                                                                                                                                                                                                                                                                  Homo sapiens.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         standard; cDNA; 149671
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       AGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCT 27203
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                                                                                                                                                                                  2000US-0237189P
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Score 45;
Pred. No.
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Detecting granulocyte activation by detecting differential expression of genes associated with granulocyte activation, which serves as diagnostic markers that is useful for monitoring disease states and drug toxicity.

WPI; 2002-435328/46.

Beazer-Barclay Y, Weissman

MS,

Yamaga Ś

Vockley

(GENB-) GENB LOGIC INC.

Claim

1; SEQ ID NO 1368; 114pp; English.

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RESULT 183
ADB70361/c
ID ADB703
XX ADB703
XX ADB703
AC ADB703
XX Moesin
XX Moesin
XX Gancer
KW Squamo
KW Homo s
XW human;
XX Homo s
XX Homo s
XX PN WO2003
XX PD 13-MAR
XX PF 05-SEF
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Best Local :
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                                                                                                                                                                                                                                                             cancer; malignant pleural mesothelioma; MPM; lung adenocarcinoma;
squamous carcinoma; medulloblastoma; prostate cancer; breast cancer;
                           05-SEP-2002; 2002WO-US028203
                                                                                                                                                                                                                                            diffuse large
                                                                                                                                                                                                                                                                                                                                                                                         04-DEC-2003
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                                                                            13-MAR-2003.
                                                                                                                        WO2003021229-A2
                                                                                                                                                                       Homo sapiens
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                                                                                                                                                                                                                                                                                                                                                                                                                                     ADB70361;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ADB70361 standard; cDNA; 149671 BP
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                                                                                                                                                                                                                                                                                                                                          CDNA
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                                                                                                                                                                                                                                                                                                                                       SEQ ID
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Conservative
                                                                                                                                                                                                                                                                                                                                                                                      (first entry)
                                                                                                                                                                                                                                          B-cell lymphoma; follicular lymphoma; ovarian cancer;
                                                                                                                                                                                                                                                                                                                                       NO:53
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100.0%; Pred. No.
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2.7e-10;
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RESULT 184
ADJ37140/c
ID ADJ371
XX ADJ371
XX ADJ371
XX 22-APR
XX Human;
XW Human;
XW Lung a
XW 10ng a
XW follic
XX JUS2003
XX US2003
XX US2003
XX 27-NOV
XX PF 05-SEF
PR 05-SEF
PR 30-AUC
XX PA (BGHM
XX (ADJ371)
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Matches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         of genes that are differentially expressed in cancerous or non-cancerous conditions, determining the expression levels of the set of genes and calculating a ratio of the expression levels of the differentially expressed genes. Mi is useful for diagnosing the presence of cancer cells or non-cancer cells in a tissue sample, where the cancer is malignant pleural mesothelioma (MPM), lung adenocarcinoma, squamous carcinoma, medulloblastoma, prostate cancer, breast cancer, diffuse large B-cell lymphoma, follicular lymphoma and ovarian cancer, and for determining prognosis or outcome of a cancer patient. The ratio of expression levels of differentially expressed genes is used as an indicator of cancer type, cancer class, and/or cancer prognosis, all of which are useful for determining a course of treatment of a patient. The present sequence encodes a human protein which is used in an example from the present
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Diagnosing cancer cells in tissue sample, or determining prognosis outcome of cancer patient, by calculating ratio of expression level genes that are differentially expressed in cancer and non cancer ti
                                                          05-SEP-2001; 2001US-0317389P.
30-AUG-2002; 2002US-0407431P.
                                                                                                                                                                                                                                Homo sapiens
                                                                                                                                                                                                                                                                                   Human; malignant pleural mesothelloma; MPM; gene; ss; tumour; lung adenocarcinoma; squamous carcinoma; medulloblastoma; prostate cancer; breast cancer; diffuse large B-cell lymphoma;
                                                                                                                                                                                                                                                                                                                                                             Human malignant pleural mesothelioma (MPM) cDNA #23
                                                                                                                                                                                                                                                                                                                                                                                                    22-APR-2004
                                                                                                                                                                                                                                                                                                                                                                                                                                                                            ADJ37140 standard; cDNA; 149671 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Sequence 149671 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             the prognosis or outcome of a cancer patient. M1 involves providing a set
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The present invention describes a method (M1) for diagnous of cancer cells or non-cancer cells in a tissue sample,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Claim 67; Page 181-263; 396pp; English.
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30-AUG-2002; 2002US-00236031.
                                                                                                                   05-SEP-2002; 2002US-00236031.
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                                                                                                                                                                                             US2003219760-A1
                                                                                                                                                                                                                                                                     follicular lymphoma;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Local
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45; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              invention describes a method (MI) for diagnosing the presence plls or non-cancer cells in a tissue sample, or determining
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Conservative
                                                                                                                                                                                                                                                                                                                                                                                                    (first entry)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          45600 A;
                                                                                                                                                                                                                                                                       ovarian
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Pred. No.
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2.7e-10;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Length 149671;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          38374 T; 0 U; 0 Other;
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(BGHM) BRIGHAM & WOMENS HOSPITAL INC.

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           cc cells or non-cancer cells in a tissue sample, determining prognosis or contcome of a cancer patient, selecting a course of treatment for a course of a cancer patient, selecting a course of treatment for a cc subject having or suspected of having malignant pleural mesothelioma (MPM) and evaluating treatment of MPM comprising determining the ratio of the expression level of a set of genes differentially expressed in a cc cancer tissue. The cancer is chosen from MPM, lung adenocarcinoma, diffuse large B-cell lymphoma, follicular lymphoma and ovarian cancer. The method is useful for diagnosing MPM in a subject suspected of having the method is useful for diagnosing MPM in a subject suspected of having the expression of nucleic acid markers or comparession products in the tissue sample suspected of being cancerous from a subject and determining the expression of nucleic acid markers or the expression products in the tissue sample. This sequence this patient cd did not form part of the printed specification but was obtained in clear content of the printed specification but was obtained in electronic format directly from USPTO at sequence data for this patient contents.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Query Match
Best Local S
Matches 45
Disclosure; SEQ ID NO 879; 198pp; English
                                   Nucleic acid array useful for detecting cancer comprises two or more nucleic acid probes.
                                                                                                                                                                                                                                                                       07-APR-2005
                                                                                                                                                                                                                                                                                                         WO2005031001-A2
                                                                                                                                                                                                                                                                                                                                                                                              Diagnosis; DNA microarray; microarray; biochip; cancer; neoplasm,
                                                                                                                                                                                                                                                                                                                                                                                                                                Human cancer-associated genomic DNA #75.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                       16-JUN-2005
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 149671 BP; 45600 A; 33308 C; 32389 G; 38374 T; 0 U; 0 Other;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  determining ratio of cancer tissues.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Diagnosing the presence of cancer or non-cancer cells in tissue sample, useful for diagnosing malignant pleural mesothelioma comprises determining ratio of expression level of a set of genes expressed in
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                                                                                                                                                                                                  23-SEP-2003; 2003US-00669920.
                                                                                                                                                                                                                                  23-SEP-2004; 2004WO-US031617.
                                                                                                                                                                                                                                                                                                                                            Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                              cytostatic; gene; ds.
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                                                                                                                                                              (CHIR )
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                                                                                         2005-273395/28
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                                                                                                                           DW, Malandro MS
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           a method of diagnosing the presence of cancer
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. 2.7e-10;
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                                                                                                            CC The invention also relates to a peptide array comprising two or more considered polypeptides encoded by a CA nucleic acid sequence, a compound CC that binds to a polypeptide, an isolated antibody or its fragment which CC binds to a polypeptide, which is prepared by immunizing a host animal CC dinds to a polypeptide, which is prepared by immunizing a host animal CC fragment and collecting cells from the host expressing antipen binding CC fragment and collecting cells from the host expressing antipen binding CC fragment and collecting cells from the host expressing antibodies against the antipody and a carrier, a method of screening for anticancer activity, a CC method of detecting a CA nucleic acid, a method of diagnosing cancer, a method of tracting cancer and a method of inhibiting expression of a CA concleic acids. The antibody is useful for detecting the presence or cabeence of cancer cells in an individual which involves contacting cells from the individual with the presence of cancer cells in an individual which involves contacting cells complex correlates with the presence of cancer cells in an individual. The composition is useful for inhibiting growth of cancer cells in an individual. The invention is also useful for dagnosing in the cancer cells in an individual. The invention is also useful for dagnosing in the cancer cells in an endividual or for delivering a therapeutic agent to cancer the invention is also useful for dagnosing in the cancer cells in an individual or for delivering a complex concerning the cancer cells in the c
Sequence 171398 BP; 51304 A; 33907 C; 34019 G;
                                                                                          the invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  associated (CA) nucleic acid, comprising two or more nucleic acid
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 The invention relates to a nucleic acid array for detecting a cancer
    52168 T;
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Matches Query Match Local 3071 CAAGATTGTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACT 3115 1 Similarity
45; Conser 1.4%; Score 45; DB llarity 100.0%; Pred. No. 2. Conservative 0; Mismatches DB 14; 2.7e-10; hes 0; Length 171398; Indels 0, Gaps

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ADL13780
ID ADL13
XX ADL1
AC ADL1
XX Oste
XX O RESULT 186 06-MAY-2004 (first entry ADL13780 standard; DNA; 190117 BP

joint space narrowing; osteophyte development; joint osteoarthritis; SNP; single nucleotide polymorphism. ds; gene; osteopathic; antiinflammatory; antiarthritic; runritic; gene therapy; joint pain;

Osteoarthritis-associated polymorphic nucleotide #312.

Homo sapiens

WO2003054166-A2

19-DEC-2002; 2002WO-US041225

20-DEC-2001; 2001US-0342603P

(INCY-) INCYTE GENOMICS

WPI; 2003-559141/52.

ζ

Schafer A;

a protein Determining susceptibility of an individual to joint space narrowing, osteophyte development and/or joint pain comprises identifying whether the individual has at least one polymorphism in a polynucleotide encoding

Disclosure; SEQ ID NO 312; 297pp; English

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RESULT 187
ABD33586
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Best Local
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The invention relates to cancer-associated proteins (CAP) and the cancer-associated (CA) nucleic acids encoding them. The invention also relates to a method for treating cancers involving administering to a patient an inhibitor of CAP, and a method of screening for anticancer activity in a potential drug involving providing a cell that expresses a CA gene, contacting a tissue sample derived from a cancer cell with an anticancer drug candidate and monitoring the effect of the anticancer drug candidate on expression of the CA gene. The CAP proteins are useful for detecting cancer associated with expression of a CAP protein in a test cell sample and for screening for a bioactive agent capable of modulating the activity of a CAP protein. The CA nucleic acids are useful for diagnosing
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  The invention relates to a method of determining susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain comprising identifying whether the individual has at least one polymorphism in a polymucleotide encoding at least one of the protein listed in the specification. The methods, composition and agent are useful for modulating the susceptibility of an individual to joint space narrowing and/or osteophyte development and/or joint pain that is associated with a disease, preferably osteoarthritis. The cell line and the non-human animal are useful for screening for an agent for diagnosing an individual having susceptibility to joint space narrowing and/or osteophyte development and/or joint pain. This sequence corresponds to the polynucleotide encoding a protein listed in the specification. (Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format directly from WIPO at
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         of cancer
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human cancer-associated (CA)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     human cancer associated protein encoded within open reading ncer associated gene, useful as targets for diagnosing cancer
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muleic acid (I). Also described: (1) an expression vector comprising (I) (2) a host cell comprising (I) or the expression vector; (3) a comprising to the above encoded within an open reading comprising that binds to the above polypeptide; (6) a hybridoma that the produces the above monoclonal antibody; (7) a pharmaceutical composition comprising the above antibody and a pharmaceutical excipient; (8) a kit for detecting cancer cells, comprising the (monoclonal) antibody conscitudes comprising the (monoclonal) antibody conscitudes comprising the (monoclonal) antibody conscitudes conscituded above; (9) methods for diagnosing cancer or for detecting the presence or absence of cancer cells in an individual; (10) a method for inhibiting growth of cancer cells in an individual; (11) a method for continuous comprising the monoclonal and individual; (12) an electronic library comprising the above polynucleotide or polypeptide, or their fragments; (13) methods of screening for anticancer activity or for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Query Match
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     14-FEB-2003; 2003US-00367094.
14-MAR-2003; 2003US-003B8838.
23-SEP-2003; 2003US-00669920.
15-DEC-2003; 2003US-00737318.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           cancer; cancer associated nucleic acid; canc
cancer associated protein; CAP; cytostatic;
lymphoma; leukaemia; human; gene; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    The present invention describes an isolated cancer associated
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Claim
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          New isolated cancer-associated polynucleotides and polypeptides useful for diagnosing, preventing or treating cancers, especially lymphoma an leukemia, or in screening for agents that modulate cancer.
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CC control is indicative of Noonan syndrome; in the test subject, and cativity of a PTPN11 protein encoded by the level of expression or activity in a control subject, where an increased expression or basal cC comprising assessing the level of expression or activity of a PTPN11 CC protein in the test subject, and comparing it to the level of expression or activity in a control subject, where an increased expression or basal cC control is indicative of Noonan syndrome; (3) treating Noonan syndrome in cC a patient, comprising administering an agent that modulates the cC expression or activity of a PTPN11 protein in association with a carrier; CC (4) an isolated PTPN11 variant comprising a mutation resulting in cC increased level of PTPN11 activity; (5) an isolated cell comprising a vector comprising a mucleic acid encoding the PTPN11 variant of (4), rector comprising a mucleic acid encoding the PTPN11 variant of (4),
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                                                                                                                                                                                                           in a subject. The method comprises detecting a mutation in the protein tyrosine phosphatase 11 (PTNN1) gene in a subject, where the mutation results in increased PTPN11 expression or activity as compared to control. The human PTPN11 gene is located on chromosome 12, more specifically to 12q24. Also described: (1) a kit for diagnosing Noonan syndrome, comprising an oligonucleotide that specifically hybridises to or adjacent to a site of mutation of a PTPN11 gene that results in increased activity of a PTPN11 protein encoded by the gene or an antibody that results fingly.
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                                                                                                                                                                                                                                                                                                                               The present invention describes a method for diagnosing Noonan syndrome in a subject. The method comprises detecting a mutation in the protein
                                                                                                                                                                                                                                                                                                                                                                              Claim 24; SEQ ID NO 33; 262pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                            Diagnosing and treating Noonan a protein tyrosine phosphatase
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          01-OCT-2001; 2001US-0326532P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          01-OCT-2002; 2002WO-US031290
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      variant; mutation;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Human PTPN11
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Noonan syndrome; protein tyrosine phosphatase 11; PTPN11; mutant;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            MOUNT SINAI SCHOOL MEDICINE
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nilarity 100.0%;
Conservative 0
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26136
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Pred.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        12;
                                                                                                                                                                                                                                                                                                                                                                                                                            syndrome in a subject using a mutation 11 gene with increased expression or
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    ВP
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      enzyme; gene;
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2.6e-10;
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/number= 4 exon 198677198793	intron 197501. 198676 /*tag= i	⋾.	/*tag= g /number=	/*Tag= r /number= 3 intron 194626197307	. 4	no .	/ rumber 2	/*tag= c /number= 136831		t= "PTPN11"	· H	key Location/Qualifiers exon 123211123604 +tag= a	thetic.	HOMO continue	<pre>lung cancer; colorectal cancer; pancreatic cancer; bladder cancer; kidney cancer; thyroid cancer; melanoma; leukaemia; human; chromosome 12; qene; ds.</pre>	heukaemia; ondition;	<pre>protein tyrosine phosphatase gene 11 variant; PTPN11 variant; haematologic disorder; mutation; increased PTPN11 activity; cytostatic; neuroprotective; PTPN11 modulator; acute lymphoblastic leukaemia; ALL;</pre>	tyrosine phosphatase gene 11; PTPN11; enzyme	in tyrosi	12-AUG-2004 (first entry)	ADO14076;	RESULT 191 ADO14076 ID ADO14076 standard; DNA; 300001 BP.		3078 GTGCCACTGCACCTCGGGCAACAGAGCAAGACTCTGTCTC 3122	Utery Match 1.4*; SCORE 45; DB 10; Length 300000; Best Local Similarity 100.0*; Pred. No. 2.6e-10; Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	300000 BP; 84671 A; 64420 C; 64260 G; 8	Aberrant expression and/or activity of the Firnit gene, specifically Noonan syndrome. The present sequence represents human PTPN11 genomic DNA, which is given in the exemplification of the present invention.	
PD 21-MAY-2004. XX	PN WO2004041216-A2	ST PT	FT exon	FT intron	FT exon		FT intron	FT exon	FT intron	FT exon		FT intron	FT exon	FT intron	FT exon	PT	FT FT intron	FT exon	FT intron		FT exon	FT FT intron FT	FT exon	FT intron	FT exon	FT intron	FT exon	FT intron
	-A2.	/*tag= af /number= 16	∵7,	/number= 15 2489102499: /*tag= ae	ρ.	⊪i po⊸	/"tag- ab /number= 14 24637024	. 13	ρı +	и.	Y T	/*cag= x /number= 12 23262423:	. "	٠.	2305882307 /*tag= v /number= 11	۳,	/*tag= t /number= 10 2263192305	. "	. 22	/*tag= r /number= 9	. "	<u>,</u> ը՝ դ	/*tag= p	٥.	/number= 6 2170572171 /*tag= n /number= 7	9 .	μ.	X. U.
			15 250510	9937	248909 d		14 248807	6369	246256 a	3288		12 233136	11 232623	2555	0742		0587	9 226318	6186		8 222128	8 221969	221843	221763	7153	217056	200176	5 200062

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RESULT 192
ADJ12734
ID ADJ127
XX
AC ADJ127
XX
DT 20-MAY
XX
DT 20-MAY
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DE DNA fr
                                                                                                                                                                                                                                                                                                                                                                                     CC comprising a mutation resulting in an increased level of prpN11 activity, where the mutation corresponds to an amino acid substitution selected CC from Asm58Tyr, Gly60Val, Asp61Tyr, Asp61Val, Glu69Iys, Phe7ILys, CC Gl176Gly, Glu76Val, Asp61Tyr, Asp61Val, Glu76Val, Glu76Val, Glu76Val, Glu76Val, Glu76Val, Glu76Val, Glu76Gly, Glu76Val, CC Gl176Gly, Glu76Val, Pro491Ser, Pro491Leu, Ser502Pro, Gly50Arg, CC Gl176Gly, Glu76Val, Pro491Ser, Pro491Leu, Ser502Pro, Gly50Arg, CC Gly503Ala, Thr507Lys, Gln510Lys, and combinations of them, in the human CC Gly503Ala, Thr507Lys, Gln510Lys, and combinations of them, in the human CC Gly503Ala, Thr507Lys, Gln510Lys, and combinations of them, in the human CC Gly603Ala, Thr507Lys, Gln510Lys, and combinations of them, in the human CC Gly603Ala, Thr507Lys, Gln510Lys, and combinations of them, in the human CC Gly703Ala, Pro491Ser, Glu76Gly, Also described: C1) characterising a mutation in the FTPN11 gene in the subject, which comprising a rector having (I) a haematologic disorder in a grant comparing it to a control; (2) a kit (II) for diagnosing a mutation results in an increased expression control sequence; (5) an isolated mucleic acid encoding (I); C2 and (6) characterising a vector having (I), operatively associated with an C2 cused as a modulator of FTPN11 protein and a carrier; (4) an isolated control. (I) has cytostatic and neuroprotective activities, and can be C2 cused as a modulator of FTPN11 activity. (W2) is useful for treating a C3 haematologic disorder such as acute lymphoblastic leukaemia (ALL), acute mysloid leukaemia (AML), in a patient. (M3) is useful for resting a cancer of pre-cancerous condition in a subject, where c the cancer is lung cancer, colorectal cancer, melanoma and leukaemia. The colorated on chromosome 12. more specifically to 12624.
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                                                                                                                                                                                                                                                                                                     Matches
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Best Local :
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                characterizing cancerous and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   05-NOV-2002; 2002US-0424170P.
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  DNA fragment of a BAC clone that encodes a human secreted protein Seq588
                                         20-MAY-2004
                                                                               ADJ12734;
                                                                                                                    ADJ12734 standard; DNA; 116
                                                                                                                                                                                                                                                                                                                                                                                   Sequence 300001 BP;
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UNIVERSITAETSKLINIKUM FREIBURG.
                                                                                                                                                                                                                                                                                                                          Similarity
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                                                                                                                                                                                                                                                 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
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                                                                                                                                                                                                                                                                                                                                                                                                                          chromosome 12,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            invention describes an isolated protein tyrosine phosphatase PN11) variant (I) associated with haematologic disorders, and
                                                                                                                                                                                                                                                                                                     Conservative
                                     (first entry)
                                                                                                                                                                                                                                                                                                                                                                                   84672 A; 64420 C; 64260 G;
                                                                                                                                                                                                                                                                                                                        1.4%; Score 45;
100.0%; Pred. No.
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                                                                                                                                                                                                                                                                                                     Mismatches
                                                                                                                                                                                                                                                                                                                                           DB 12;
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                                                                                                                                                                                                                                                                                                                                                                                   85849 T; 0 U; 800 Other;
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This invention relates to novel polynucleotides encoding human secreted proteins. Specifically, it refers to the vectors, host cells, recombinant and synthetic methods for producing human polynucleotides, polypeptides and antibodies. Furthermore, it relates to screening methods to identify agonists and antagonists that can be used to inhibit or enhance the production and function of the secreted proteins. The present invention describes these compositions as useful for diagnosing, treating or preventing disorders such as cancer, haematopoietic diseases including prostatitis and inguinal hermia, musculoskeletal diseases including systemic lupus crythematosus and gout, cardiovascular disease including systemic lupus crythematosus and gout, cardiovascular disease including

arrhythmia and hypernatraemia,

mixed

New isolated nucleic acids and polypeptides, useful for diagnosing, treating, preventing or ameliorating diseases or disorders e.g. cancer, anemia, arthritis, asthma, inflammatory bowel disease or Alzheimer's

Disclosure;

SEQ

ID NO 588; 286pp; English

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09-OCT-1997;
09-OCT-1997;
09-OCT-1997;
09-OCT-1997;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 multiple myeloma; reproductive system disorder; prostatitis; myeloma; reproductive system disorder; prostatitis; inguinal hernia; musculoskeletal disease; systemic lupus erythematosus; gout; cardiovascular disease; arrhythmia; hypernatraemia; fetal disease; urinary incontinence; renal disorder; neural; sensory disease; alzheimer's disease; meningitis; respiratory disease; emphysema; occupational lung disease; endocrine disease; diabetes; glomerulonephritis; digestive disease; portal hypertension; irritable bowel syndrome; epithelial disease; scleroderma; epidermolysis bullosa; cytostatic; antianemic; antiarthritic; antiasthmatic; anti-AITV; immunosuppressive; antisirhamatory; antiasthria; antibacterial; osteopathic; dermatological; antigout; nephrotropic; uropathic; neuroprotective; antiparkinsonian; tranquilizer; nephrotropic; uropathic; neuroprotective; antiparkinsonian; tranquilizer;
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(FLOR/)
(GREE/)
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                                                              OLSEN H.
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                                                                                    BBNER
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Brewer LA,
Ferrie AM,
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97US-0061527P.
97US-0061529P.
97US-0061532P.
97US-0061536P.
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Duan RD, Ruben SM,
Yu G, Florence C,
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Florence KA, Gre
Ebner R, Olsen H;
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                      Greene JM;
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RESULT 193
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Best Local S
Matches 44
 31-JAN-2000

04-FEB-2000

24-FEB-2000

24-FEB-2000

12-MAR-2000

11-MAR-2000

11-MAR-2000

11-MAY-2000

07-JUN-2000

28-JUN-2000

07-JUL-2000

07-JUL-2000

11-JUL-2000

11-JUL-2000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             alcohol syndrome and Down's syndrome, excretory diseases including urinary incontinence and renal disorders, neural or sensory disease including Alzheimer's disease and meningitis, respiratory disease including emphysema and occupational lung disease, endocrine diseases including diabetes and glomerulonephritis, digestive diseases including portal hypertension and irritable bowel syndrome and connective tissue of epithelial diseases including soleroderma and epidermolysis bullosa. As such, there are various activites such as cytostatic, antianemic, antiarthritic, antiasthmatic, anti-HIV, immunosuppressive,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                     17-JAN-2001; 2001WO-US001354.
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                                                                                                                                                                                                                                                                                                                                                                                                                                      WO200157182-A2
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human immune/haematopoietic antigen genomic sequence SEQ ID NO:22193.
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2000US-0184664P.
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2000US-0199076P.
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2000US-0199123P.
2000US-0214886P.
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2000US-0216647P.
2000US-021669P.
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2000US-0224518P.
2000US-0225214P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       DB 12; Length 116; 1e-09;
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 2000US-0225447P.
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                                             CC AAKS4951 to AAK64702 encode the human immune/haematopoietic antigen (I) CC amino acid sequences given in AAM62170 to AAM91921. (I) have cytostatic CC activity, and can be used in gene therapy and vaccine production. (I) CC proteins and polynucleotides may be used in the prevention, diagnosis and CC treatment of diseases associated with inappropriate (I) expression. For CC example, they may be used to treat disorders associated with decreased CC expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to CC supplement the patients own production of (I). Additionally, (I) CC polynucleotides may be used to produce the secreted (I), by inserting the CC protein. (I) proteins and polynucleotides may be used to prevent, CC diagnose and treat immune/haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoietic actigen genomic CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169 crepresent sequences used in the exemplification of the present invention
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AAK7444
AC AAK744
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Human;
KW Cytost
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U02-010
PR 24-FEB
PR 04-FEB
PR 11-JUL
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Matches 44
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26-JUL-2000

14-AUG-2000

15-SEP-2000

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01-SEP-2000
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07-JUL-2000;
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19-MAY-2000;
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28-JUN-2000;
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  2000US-0186350P

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2000US-0231244P,
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2000US-0234274P,
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2000US-024677P,
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Best Local Similarity
Matches 44; Conserva
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Human; 5' \mathtt{EST}_i expressed sequence tag; secreted protein; cDNA isolation; gene therapy; chromosome mapping; ss.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.
                                                              Human secreted protein 5' EST, SEQ ID NO: 25910
                                                                                                        06-OCT-2000 (first entry)
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                                                                                                                                                                                     AAC21835 standard; cDNA; 145
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        29269; 3071pp + Sequence Listing; English
                                                                                                                                                                                                                                                                                                                                                                         0;
                                                                                                                                                                                                                                                                                                                                                                                          Score 44; DB 4;
Pred. No. 1e-09;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        MS.
                                                                                                                                                                                     ВP
                                                                                                                                                                                                                                                                                                                                                                         Mismatches
                                                                                                                                                                                                                                                                                                                                                                         0;
                                                                                                                                                                                                                                                                                                                                                                                                             Length 139;
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Gaps

0

06-SEP-2000
08-SEP-2000
08-SEP-2000
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08-SEP-2000
08-SEP-2000
08-SEP-2000
08-SEP-2000
08-SEP-2000
14-SEP-2000
14-SEP-2000
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14-SEP-2000
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25-SEP-2000
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29-SEP-2000
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29-SEP-2000
29-SEP-2000
20-OCT-2000
20-OCT-2000
20-OCT-2000
02-OCT-2000
02-OCT-2000
02-OCT-2000
02-OCT-2000
02-OCT-2000
03-NOV-2000
03-NOV-2000
03-NOV-2000
04-NOV-2000
06-NOV-2000
08-NOV-2000
08-NOV-2000
01-NOV-2000

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RESULT 196
AAK91189
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Query Match
Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      The present sequence is one of a large number of 5' ESTs derived from mRNAs encoding secreted proteins. No ORF has yet been conclusively identified within the present sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of mRNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design
                                                                                                                                                                                                 Human; digestive system antigen; gene therapy; cancer; ulcerative colitis; infection; Hirschsprung's disease; digestive system disorder; Meckel's diverticulum; ds.
 31-JAN-2000; 2000US-0179065P.
04-FEB-2000; 2000US-0180628P.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     New nucleic acid that is a 5' expressed sequence tag (5' EST) for obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Dumas Milne Edwards J,
                                                      17-JAN-2001; 2001WO-US001324.
                                                                                                                                                                                                                                                                         Human digestive system antigen genomic sequence SEQ ID NO: 4765.
                                                                                                                                                                                                                                                                                                                05-NOV-2001
                                                                                                                                                                                                                                                                                                                                                  AAK91189;
                                                                                                                                                                                                                                                                                                                                                                                    AAK91189 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Sequence 145 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           expression and secretion vectors
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Claim 1; SEQ ID NO 25910; 71pp + Sequence Listing; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    diagnostic, forensic, gene therapy and chromosome mapping procedures
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       26-FEB-1999;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           21-FEB-2000; 2000EP-00200610.
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                                                                                            02-AUG-2001.
                                                                                                                             WO200155314-A2
                                                                                                                                                                 Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     (GEST ) GENSET.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   3079 TGCCACTGCACTCCAGCCTGGGCAACAGAGCCAAGACTCTGTCTC 3122
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      44;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                TGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Conservative
                                                                                                                                                                                                                                                                                                              (first
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         99US-0122487P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         25 A; 48 C; 38 G;
                                                                                                                                                                                                                                                                                                                                                                                    DNA;
                                                                                                                                                                                                                                                                                                              entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1.4%;
                                                                                                                                                                                                                                                                                                                                                                                      182
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Score 44;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                      ₽₽
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Giordano
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Ţ,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          DB 3; L. J. 1e-09; 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         0 U;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 <u>.</u>
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         2 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Length 145;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                4
                                                                                                                                                                                                                        appendicitis;
chronic colitis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Gaps
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0
   14-AUG-2000
12-AUG-2000
12-AUG-2000
13-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
11-AUG-2000
11-AUG
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02-MAR-2000;
16-MAR-2000;
17-MAR-2000;
18-APR-2000;
19-MAY-2000;
07-JUN-2000;
28-JUN-2000;
30-JUN-2000;
07-JUL-2000;
07-JUL-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              11-JUL-2000;
14-JUL-2000;
26-JUL-2000;
2000US-020515P

2000US-0214866P

2000US-0216860P

2000US-0216860P

2000US-0216860P

2000US-021680P

2000US-0218290P

2000US-02254519P

2000US-02252119P

2000US-02252119P

2000US-0225266P

2000US-0225268P

2000US-0225268P

2000US-0225759P

2000US-0225759P

2000US-02268681P

2000US-02268681P

2000US-02271829P

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2000US-0233140P

2000US-02331413P

2000US-0233143P

2000US-0233143P

2000US-0233143P

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2000US-0233339P

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2000US-0233339P

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2000US-02333339P

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2000US-0186350P.
2000US-0189874P.
2000US-0190076P.
2000US-0198123P.
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02-OCT-20001; 2000015-02370339; CX 2000015-02370339; CX 2000015-02370319; CX 2000015-02370319
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31-JAN-2000 04-FBB 2000 24-FBB 2000 16-MAR-2000 11-MAR-2000 11-MAR-2000 11-MAR-2000 11-MAY-2000 07-JUN-2000 07-JUN-2000 07-JUL-2000 07-JUL-2000 11-JUL-2000 11-JUL-2000

2000US-0189874P.
2000US-0190176P.
2000US-0198123P.
2000US-0205515P.
2000US-020148867P.
2000US-021148867P.
2000US-02114887P.
2000US-02116847P.
2000US-0217487P.
2000US-0217487P.
2000US-0217487P.
2000US-0217496P.
2000US-02182996P.
2000US-022963P.
2000US-0228518P.
2000US-0224518P.
2000US-0225213P.
2000US-0225213P.
2000US-0225213P.
2000US-0225214P.
2000US-0225214P.

WO200155355-A1 02-AUG-2001. 17-JAN-2001; 20

2001WO-US001351. 2000US-0179065P. 2000US-0180628P. 2000US-0184664P. 2000US-0186350P. sapiens.

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RESULT 197
AAS32121
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Query Match
Best Local S
Matches 44
cerebrovascular disorder; nervous system disorder; bacterial infection; fungal infection, viral infection; ocular disorder; endocrine disorder; gastrointestinal disorder; renal disorder; respiratory disorder; wound healing; skin aging; organ transplantation; tissue regeneration; anti-infertility.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The present invention provides the protein and coding sequences of number of human digestive system antigens. These can be used in the diagnosis, treatment and prevention of diagnosis, treatment and prevention on the system disorders, including cancer, Meckel's diverticulum, bacterial or parasitic infections, appendicitis, Hirschsprung's disease, chronic colitis ulcerative colitis. The present sequence is a genomic DNA fragment
                                                                                                                  Liver associated protein; human; mouse; rabbit; goat; horse; cat; dog; chicken; sheep; immunosuppregsive; antiarthritic; vasotropic; antiproliferative; cytostatic; cardiant; neuroprotective; cerebroprotective; nootropic; antibacterial; virucide; fungicide; cancer. ophthalmological; vulnerary; gene therapy; autoimmune disease; neoplasm; hyperproliferative disorder; breast; liver; cardiovascular disorder; ds;
                                                                                                                                                                                                                                                                                       Human liver associated genomic DNA #295.
                                                                                                                                                                                                                                                                                                                                     04-DEC-2001
                                                                                                                                                                                                                                                                                                                                                                                   AAS32121;
                                                                                                                                                                                                                                                                                                                                                                                                                               AAS32121
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 182 BP; 48
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGAGCTCTGTCT 3121
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 139 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 182
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              44;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      a digestive system antigen of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                               standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Conservative
                                                                                                                                                                                                                                                                                                                                     (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 1.4%; Sur
, 100.0%; Pr
                                                                                                                                                                                                                                                                                                                                                                                                                               DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     A; 44 C;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Score 44;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       55 G; 35 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Mismatches
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le-09;
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Sequences AAS31827-AAS32182 represent genomic DNA molecules, which encode the liver associated polypeptides of the invention. Liver associated CC polypeptides and their associated polypuclectides are useful in the CC diagnosis, treatment and prevention of various types of disorders in e.g. CC humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. A pathological condition can be determined by detecting the presence or absence of a mutation in a liver associated polynucleotide. The treatable CC disorders include autoimmune diseases such as rheumatoid arthritis, CC hyperproliferative disorders such as neoplasms of the breast or liver, CC disorders for a such as cardiac arrest, cerebrovascular disorders such as cardiac arrest, cerebrovascular CC disorders such as cerebral ischaemia, nervous system disorders such as corneal infection, endocrine disorders such as CC premature labour and infertility, gastrointestinal disorders such as CC crohn's disease, renal disorders such as glomerulonephritis and CC respiratory disorders such as asthma and pleurisy. The polypeptides can calso be used to aid wound healing, to prevent skin aging due to sunburn, to maintain organs before transplantation, to regenerate tissues and in chemotaxis. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly
            08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
08-NOV-2000
17-NOV-2000
17-NOV
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Isolated nucleic acid used in preventing, to particularly cancer of
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Claim
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    (HUMA-)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             2000US-0246526P

2000US-0246528P

2000US-0246532P

2000US-0246610P

2000US-0246611P

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2000US-0249210P

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2000US-0249215P

2000US-0249216P

2000US-0249217P

2000US-0251865P

2000US-0251866P

2000US-0251866P

2000US-0251866P

2000US-0251869P

2000US-0251869P

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2000US-0251989P

2000US-0251989P

2000US-0251990P

2001US-025199P

2001US-025199P
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treating or a
r of the liver.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  526pp;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        encoding a human liver related rameliorating disorders of the
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14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
14-AUG-2000
22-AUG-2000
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23-SEP-2000
23-SEP-2000
24-SEP-2000
25-SEP-2000
26-SEP-2000
27-SEP-2000
28-NOV-2000
29-SEP-2000
20-OCT-2000
20-OCT

2000US.

S-0229345P.
S-02295199P.
S-0239437P.
S-0230438P.
S-0231242P.
S-0231244P.
S-0231244P.
S-0231244P.
S-0231413P.

S-0226681P.
S-0226868P.
S-02271082P.
S-0227092P.
S-0228924P.
S-02289287P.
S-0229343P.
S-0229344P.

S-0225447P. S-0225757P. S-0225758P. S-0225759P. S-0226279P.

S-0232080P S-0231968P S-0231968P S-023239PP S-02323064P S-023400P S-023492PP S-023492PP S-023492PP S-023493064P S-023499PP S-023484P S-023486PP S-023686PP S-023686PP S-023687PP S-023687PP S-023687PP S-0237039PP S-0237039PP S-0237039PP S-0237039PP S-0237039PP S-0237039PP S-0241786PP S-02446477PP S-02446477PP

protein

8

from

WIPO at ftp.wipo.int/pub/published_pct_sequences

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RESULT 198
ABN90476
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Query Match
 14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
14-AUG-2000;
22-AUG-2000;
30-AUG-2000;
01-SEP-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human; liver antigen; liver disorder; hepatic disorder; infection; hepatitis; viral; parasitic; bacterial; fungal; inflammatory condition; cirrhosis; granulomatous hepatitis; toxin damage; drug damage; autoimmune disease; Wilson's disease; primary biliary cirrhosis; neoplastic disorder; cancer; tumour; portal hypertension; gastrointestinal disorder; hepatitis; drug screening; gene therapy; chromosome mapping; forensic analysis; antibody preparation; hepatotropic; cytostatic; antiinflammatory; virucide; antibacterial; fungicide; parasiticide; antidote; immunosuppressive; gene; ds.
                                                                                                                                                                                                                                                                                                                                                                     31-JAN-2000;
04-FEB-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                11-APR-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                         US2002042096-A1
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human liver antigen HHLAB49 genomic sequence, SEQ ID NO:597
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       24-JUL-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ABN90476;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ABN90476 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 182 BP; 48 A; 44 C; 55 G; 35 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                         17-JAN-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 3121
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                GTGCCACTGCAGCCTGGGCAACAGAGCAAGACTCTGTCT 182
2000US-0214886P.
2000US-021680P.
2000US-021680P.
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2000US-0224518P.
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2000US-0225266P.
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2000US-0225757P.
2000US-0225789P.
2000US-0225789P.
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2000US-022989P.
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2000US-022989P.
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2000US-023421P.
2000US-023421P.
2000US-023421P.
2000US-023427P.
2000US-023427P.
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2000US-023427P.
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2000US-0180628P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 0; Mismatches
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1e-09;
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CC encompasses polypeptides 90% identical and polynucleotides 95% identical control of the sequences of the invention. The invention additionally relates to combinant vectors and host cells comprising human liver antigen composing or preventing various disorders of the liver. Such conditions (c) polynucleotides, antibodies against human liver antigens, and the use of collever antigen polynucleotides and polypeptides in diagnosing, treating, composing or preventing various disorders of the liver. Such conditions (c) include viral infections (e.g., cytomegalovirus, Espetin-Barr virus, hepatitis A virus, hepatitis B virus and hepatitis C virus), parasitic cinfections (e.g., Clonorchis sinensis, Echinococcus granulosus and confections (e.g., Confections, Confections,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         29-SBP-2000
29-SBP-2000
29-SBP-2000
29-SBP-2000
02-OCT-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               The invention relates to 145 novel human liver antigens (ABP40831-ABP40975) and to cDNAs encoding them (ABN90036-ABN90180), and also
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Disclosure; SEQ ID NO 597; 181pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  New nucleic acid encoding human liver antigens, useful for diagnosis, treatment and prevention of e.g. hepatitis and hepatic cancer, also
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      WPI; 2002-381944/41.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           related polypeptides and antibodies.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       08-DEC-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           17-NOV-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ROSEN
RUBEN
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              BARASH S
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2000US-023636BP.
2000US-023636PP.
2000US-0236370P.
2000US-0237037PP.
2000US-0237038P.
2000US-0237039P.
2000US-0237039P.
2000US-023704PP.
2000US-024785PP.
2000US-0241785PP.
2000US-0241809P.
2000US-0244817P.
2000US-02494617P.
2000US-02494617P.
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2000US-02494617P.
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Sequence 182 BP; 48 Ą, 44 C; 55 G; 35 T; 0 U; 0 Other;

6,

Length 182

0

δ 뭉 Query Match Best Local S Matches Local Similarity 3078 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGAGCTCTGTCT 3121 139 GTGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCT 182 44; Conservative 1.4%; Score 44; DB; Pred. No. 1e-0; Mismatches 0; 1e-09; <u>,</u> Indels 0, Gaps

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RESULT 199
ADJ15389
IID ADJ153
XX
ADJ15389
AC ADJ153
XX
XX
IIVER
XX
IIVER
X
31-JAN-2000
04-FBB-2000
24-FBB-2000
12-MAR-2000
11-MAR-2000
11-MAR-2000
11-MAR-2000
07-JUL-2000
01-AUG-2000
01-SEP-2000
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    liver; virucide; fungicide; antibacterial; antiparasitic; hepatotropic; antiinflammatory; cytostatic; litholytic; antirheumatic; antiarthritic; neuroprotective; antidabetic; anticoagulant; thrombolytic; antiarteriosclerotic; cardiant; haemostatic; antiarrhythmic; ophthalmological; antiarteriosclerotic; vasotropic; osteopathic; nootropic; antiparkinsonian; anticonvulsant; neuroleptic; vasotropic; prostatic; gynaecological; viral; fungal; bacterial; cytostatic; gynaecological; viral; fungal; bacterial; parasitic infection; cirrhosis; Wilson's disease; gastrointestinal disorder; pancreatic; gallbladder; immune; blood; hyperproliferative; cardiovascular; respiratory; musculoskeletal system; neurological; endocrine; reproductive system; developmental; inherited; neurological; endocrine; reproductive system; developmental; inherited;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       14-FEB-2002;
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2000US-0179065P.
2000US-0184664P.
2000US-0184664P.
2000US-0184664P.
2000US-0199076P.
2000US-0199076P.
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2000US-021643P.
2000US-0217486P.
2000US-0217486P.
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2000US-0225214P.
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2000US-0225276P.
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2000US-0225759P.
2000US-0225758P.
2000US-0225758P.
2000US-0225758P.
2000US-0225759P.
2000US-0225934P.
2000US-0225934P.
2000US-0225934P.
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 S-0214886P.
S-021664PP.
S-0216880P.
S-0217496P.
S-022963P.
S-0224513P.
S-0225214P.
S-0225266P.
S-0225266P.
S-0225270P.
S-0225759P.
S-0225719P.
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25-SEP-2000
25-SEP
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RESULT 200
AAK87671/c
ID AAK876
XX
AC AAK876
XX
AC AAK876
XX
OT O7-NOV
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                                                                                                                                                                                                                                                                                                                                                                      The invention relates to a novel isolated, liver related polypeptide. The CC polypeptide of the invention demonstrates virucide, fungicide, complypeptide of the invention demonstrates virucide, fungicide, complypeptide of the invention demonstrates virucide, fungicide, complypeptide of the invention, antiinflammatory, cytostatic, complypeptide, antiinflammatory, cytostatic, antiinflammatory, antiidiabetic, antiincomplytic, antiinreteriosclerotic, cardiant, haemostatic, complypeptide, complypeptide, complypeptides, comply
                                                                                                                                                                                                                                                                          Query Match
Best Local Similarity
Matches 44; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              17-NOV-2000;
17-NOV-2000;
17-NOV-2000;
01-DEC-2000;
01-DEC-2000;
05-DEC-2000;
05-DEC-2000;
06-DEC-2000;
06-DEC-2000;
08-DEC-2000;
08-DEC-2000;
08-DEC-2000;
08-DEC-2000;
08-DEC-2000;
08-DEC-2000;
  07-NOV-2001
                                            AAK87671;
                                                                                  AAK87671 standard;
                                                                                                                                                                                                                                                                                                                                                                                   the specification per se web-site.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 New liver related polypeptide, useful for diagnosis, treatment and/or prevention of liver, gastrointestinal, pancreatic, immune, blood related, endocrine, reproductive, hyperproliferative or reproductive disorders.
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7-NOV-2000;
7-NOV-2000;
7-NOV-2000;
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0; 2000US-024924SP.
0; 2000US-024926AP.
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0; 2000US-02517198BP.
0; 2000US-0251719BBP.
0; 2000US-025186BP.
0; 2000US-025198P.
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  (first entry)
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                                                                                    DNA;
                                                                                                                                                                                                                                                                                              1.4%;
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                                                                                    307
                                                                                    ВP
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0; Mismatches
                                                                                                                                                                                                                                                                             0,
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1e-09;
                                                                                                                                                                                                                                                                          0;
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Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                              Human
                                                                                                          immune/haematopoietic antigen genomic sequence
                                                                                                              SEQ ID NO:42483
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Homo sapiens.

WO200157182-A2

17-JAN-2001; 2001WO-US001354

2000US-0179065P

17-MAR-2000; 18-APR-2000; 2000US-0184664P.
2000US-0189874P.
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2000US-023575P

19-MAY-2000 07-JUN-2000 28-JUN-2000 30-JUN-2000 07-JUL-2000 07-JUL-2000 11-JUL-2000 11-JUL-2000 14-JUL-2000 26-JUL-2000 26-JUL-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000

14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 14-AUG-2000 12-AUG-2000 22-AUG-2000 23-AUG-2000 23-AUG-2000 30-AUG-2000 01-SEP-2000 01-SEP-2000

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14-SEP-2000
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2000US-0246477P
2000US-0246610P
2000US-0246611P
2000US-024661P
2000US-0249211P
2000US-0259391P
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5
                                                                                                                                                             cc amino acid sequences given in AAM82170 to AAM91921. (1) have cytostatic cc activity, and can be used in gene therapy and vaccine production. (1) cc proteins and polynucleotides may be used in the prevention, diagnosis and ct treatment of diseases associated with inappropriate (1) expression. For cc example, they may be used to treat disorders associated with decreased cc expression by rectifying mutations or deletions in a patient's genome ct that affect the activity of (1) by expressing inactive proteins or to cc supplement the patients own production of (1). Additionally, (1) cc polynucleotides may be used to produce the secreted (1), by inserting the cc protein. (1) proteins and polynucleotides may be used to prevent, (2) diagnose and treat immune/haematopoietic related diseases, especially cc cancers and cancer metastases of haematopoietic diseases, especially cc cancers and cancer metastases of haematopoietic acitigen genomic cc sequences from the present invention. AAK54942 to AAK54950 and AAM82169 cc represent sequences used in the exemplification of the present invention.
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                                                                  Query Match
Best Local Similarity
Matches 44; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        08-DEC-2000; 2000US-0251999P
08-DEC-2000; 2000US-0251999P
11-DEC-2000; 2000US-0254097P
05-JAN-2001; 2001US-0259678P
                                                                                                                                      Sequence 307 BP; 53
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Nucleic acids encoding
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Disclosure;
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74
                       TGCCACTGCACTCCAGCCTGGGCAACAGAGCAAGACTCTGTCTC 3122
 TGCCACTGCAGCCTGGGCAACAGAGCAAGACTCTGTCTC
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                                                                1.4%; So ilarity 100.0%; I Conservative 0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                  encode the human immune/haematopoietic antigen given in AAM82170 to AAM91921. (I) have cytosts
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                                                                  Score 44; DB Pred. No. 9.8 0; Mismatches
                                                                                                                                      69 G;
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                                                            DB 4;
J. 9.8e-10;
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                                                                     Indels
   31
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polypeptides, and metastasis.

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and

the

Search completed: May Job time: 1822 secs 11, 2006, 04:27:01

0

Gaps

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